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E12
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1445930 NTIS Accession Number: PB89-200315
  Novel Interleukin 2 Receptor and Applications Thereof
  (Patent Applicatio)
  Waldmann, T. A.; Leonard, W. J.
  Department of Health and Human Services, Washington, DC.
  Corp. Source Codes: 068119000
  Report No.: PAT-APPL-7-165 302
  Filed 8 Mar 88
                  18p
  Languages: English Document Type: Patent
  Journal Announcement: GRAI8916
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1351659 NTIS Accession Number: PB88-140231
  Method for Treating Maligancy and Autoimmune Disorders in Humans
  (Patent Applicatio)
  Waldmann, T. A.
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E2 E3

E4

E5

E9

Department of Health and Human Services, Washington, DC. Corp. Source Codes: 068119000 Report No.: PAT-APPL-7-085 707 Filed 17 Aug 87 18p Languages: English Document Type: Patent Journal Announcement: GRAI8808 Government-owned invention available for U.S. licensing and, for foreign licensing. Copy of application available NTIS. Order possibly, this product from NTIS by: phone at 1-800-553-NTIS (U.S. customers); (703)605-6000 (other countries); fax at (703)321-8547; and email at NTIS is located at 5285 Port Royal Road, orders@ntis.fedworld.gov. Springfield, VA, 22161, USA. NTIS Prices: PC A03/MF A01 1/3/3 DIALOG(R)File 6:NTIS Comp&distr 1998 NTIS, Intl Copyright All Righ. All rts. reserv. 1325945 NTIS Accession Number: PB87-224424 Novel Interleukin 2 Receptor and Applications Thereof (Patent Application) Waldmann, T. A. Department of Health and Human Services, Washington, DC. Corp. Source Codes: 068119000 Report No.: PAT-APPL-7-066 989 Filed 29 Jun 87 16p Languages: English Document Type: Patent Journal Announcement: GRAI8723 Government-owned invention available for U.S. licensing and, possibly, for foreign licensing. Copy of application available NTIS. Order this product from NTIS by: phone at 1-800-553-NTIS (U.S. customers); (703)605-6000 (other countries); fax at (703)321-8547; and email at orders@ntis.fedworld.gov. NTIS is located at 5285 Port Royal Road, Springfield, VA, 22161, USA. NTIS Prices: PC A02/MF A01 1/3/4 DIALOG(R) File 6:NTIS Comp&distr 1998 NTIS, Intl Copyright All Righ. All rts. reserv. 1320089 NTIS Accession Number: PB87-218426 Novel Interleukin 2 Receptor and Applications Thereof (Patent Application) Waldmann, T. Department of Health and Human Services, Washington, DC. Corp. Source Codes: 068119000 Report No.: PAT-APPL-7-066 989 Filed 29 Jun 87 18p Languages: English Document Type: Patent Journal Announcement: GRAI8721 Government-owned invention available for U.S. licensing and, possibly, for foreign licensing. Copy of application available NTIS. NTIS Prices: Not available NTIS ? begin medtext 15mar00 08:06:04 User208760 Session D1481.3 0.223 DialUnits File6 \$1.31 \$7.20 4 Type(s) in Format 3 \$7.20 4 Types Estimated cost File6 \$8.51 \$0.10 TYMNET \$8.61 Estimated cost this search

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         (c) 2000 Mass. Med. Soc.
  File 457: The Lancet 1986-2000/Mar W1
         (c) 2000 The Lancet, Ltd.
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           32314 MG
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00107152
COPYRIGHT American Medical Association 1998
Cutaneous Graft-versus-Host Disease (ARTICLE)
  ARACTINGI, SELIM; CHOSIDOW, OLIVIER
  Archives of Dermatology
  May, 1998; Review: tzd602
  LINE COUNT: 00989
           (Item 2 from file: 442)
DIALOG(R) File 442: AMA Journals
(c) 2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.
COPYRIGHT American Medical Association 1996
Detection of Human T-Cell Lymphotrophic Virus Type I in Archival Tissue
Specimens (ARTICLE)
                 S.; RUFFO, ALICE; SALVEKAR, ANUPAMA; HENGHOLD, WILL;
          GARY
   TAKESHITA, MORISHIGE; KIKUCHI, MASAHIRO
  Archives of Dermatology
  Nov, 1996; Study: tzd1339
  LINE COUNT: 00413
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                                                      Dermatitis Selective
                                             Atopic
Circulating
              Skin-Homing
                            Т
                                Cells in
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Up-regulation of HLA-DR, Interleukin-2R, and CD30 and Decrease After
Combined UV-A and UV-B Phototherapy (ARTICLE)
  PILETTA, PIERRE A.; WIRTH, SUSANNE; HOMMEL, LUCETTE; SAURAT, JEAN H.;
  HAUSER, CONRAD
  Archives of Dermatology
  Oct, 1996; Study: tzd1171
  LINE COUNT: 00464
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           (Item 4 from file: 442)
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00094516
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Peering at the Dermatology Literature (ARTICLE)
 Archives of Dermatology
 May, 1995; Editorials: de 602
 LINE COUNT: 00366
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00093606
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       T-Cell
               Lymphotropic Virus Type
                                             I-Associated
                                                            Adult T-Cell
Leukemia The Joseph Goldberger Clinical Investigator Lecture (ARTICLE
 WALDMANN, THOMAS A.
  JAMA, The Journal of the American Medical Association
 March 1, 1995; 9 of Health: tzj735
 LINE COUNT: 00348
           (Item 6 from file: 442)
DIALOG(R) File 442: AMA Journals
(c) 2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.
00090827
COPYRIGHT American Medical Association 1994
Recurrence of Pemphigus Vulgaris Associated With Interleukin 2 Therapy (
ARTICLE)
  PRUSSICK, RONALD; PLOTT, R. TODD; STANLEY, JOHN R.
  Archives of Dermatology
  July, 1994; Observation: de_890
 LINE COUNT: 00266
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           (Item 7 from file: 442)
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(c)2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.
00090215
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Primary Gastric T-Cell Lymphoma Morphological and Immunohistochemical

Studies of Two Cases (ARTICLE)

Archives of Pathology and Laboratory Medicine May, 1994; Original Article: p547 LINE COUNT: 00257

2/3/8 (Item 8 from file: 442)
DIALOG(R)File 442:AMA Journals
(c)2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

00087400

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(ARTICLE)

GOLDMAN, BARRY D.; OH, SE-KYUNG; DAVIS, BRET E.; KADIN, MARSHALL E.; POIESZ, BERNARD J.; KOH, HOWARD K.

Archives of Dermatology

Sep, 1993; OBSERVATION: p1166

LINE COUNT: 00294

2/3/9 (Item 9 from file: 442)
DIALOG(R)File 442:AMA Journals
(c)2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

00081309

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Use and Interpretation of Diagnostic Immunologic Laboratory Tests (ARTICLE)

LOPEZ, MANUEL; FLEISHER, THOMAS; DESHAZO, RICHARD D. JAMA, The Journal of the American Medical Association November 25, 1992; 20: p2970 LINE COUNT: 01953

2/3/10 (Item 10 from file: 442)
DIALOG(R)File 442:AMA Journals
(c)2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

00055691

Administration of a Prostaglandin Synthetase Inhibitor Associated With an Increased Immune Cell Infiltrate in Squamous Cell Carcinoma of the Head and Neck (Article)

Shikani, Alan H. MD; Richtsmeier, William J. MD, Klein, Jerry L. PhD; Kopher, Kenneth A. MS Archives of Otolaryngology-Head & Neck Surgery 1992; 118: 526 (3)

2/3/11 (Item 11 from file: 442)
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00055031

Interleukin 2 and Granulocyte-Macrophage Colony-Stimulating Factor Induce a Perivascular Lymphocytic Infiltrate in a Skin Explant Model (Article)

Horn, Thomas D., MD; Rest, Ellen B., MD; Karp, Judith E., MD; Burke, Philip J., MD; Vogelsang, Georgia B., MD; Boucher, Correne L.; Hood, Antoinette F., MD
Archives of General Psychiatry 1991; 48: 1789 (5)

2/3/12 (Item 12 from file: 442)
DIALOG(R)File 442:AMA Journals
(c)2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

00050927

Large-cell Anaplastic (Ki-1-Positive) **Lymphoma** Complicated by Disseminated Intravascular Coagulation (Article)

Arber, Daniel, MD; Bilbao, Jorge, MD; Bassion, Susan, PhD Archives of Pathology & Laboratory Medicine 1991; 115: 188 (5)

2/3/13 (Item 13 from file: 442)
DIALOG(R)File 442:AMA Journals
(c)2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

00050311

Intravenous Administration of Recombinant Human Granulocyte-Macrophage Colony-Stimulating Factor Causes a Cutaneous Eruption (Article)

Horn, Thomas D., MD; Burke, Philip J., MD; Karp, Judith E., MD; Hood,
Antoinette F., MD
Archives of Dermatology
1991; 127: 49 (4)

2/3/14 (Item 14 from file: 442)
DIALOG(R)File 442:AMA Journals
(c)2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

00049917

Human T-cell **Leukemia/Lymphoma** Viruses: Life Cycle, Pathogenicity, Epidemiology, and Diagnosis (Article)

Hjelle, Brian, MD
Archives of Pathology & Laboratory Medicine
1991; 115: 440 (11)

2/3/15 (Item 15 from file: 442)
DIALOG(R)File 442:AMA Journals
(c)2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

00046690

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Childhood Ki-1 Lymphoma; A Report of Two Cases (ORIGINAL ARTICLES)

OKA, KUNIYUKI; MORI, NAOYOSHI; KOJIMA, MIZU; IIJIMA, TATSUO; HANADA, TAKASHI; TSUCHIDA, MASAHIRO
Archives of Pathology and Laboratory Medicine
September, 1989; 113: 998-1002
LINE COUNT: 00166 WORD COUNT: 02302

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DIALOG(R)File 442:AMA Journals
(c)2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

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Cutaneous-Type Adult T-Cell **Leukemia/Lymphoma**; A Unique Clinical Feature With Monoclonal T-Cell Proliferation Detected by Southern Blot Analysis (OBSERVATIONS)

TAKAHASHI, KENZO; TANAKA, TOSHIHIRO; FUJITA, MAYUMI; HORIGUCHI, YUJI;

MIYACHI, YOSHIKI; IMAMURA, SADAO

Archives of Dermatology March, 1988; 124: 399-404

LINE COUNT: 00225 WORD COUNT: 03115

2/3/17 (Item 17 from file: 442)

DIALOG(R) File 442: AMA Journals

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00038322

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Human T-Cell Lymphotropic Virus Type I-Associated Adult T-Cell Leukemia/Lymphoma in an Atypical Host (ORIGINAL ARTICLE)

GOLDMAN-LEIKIN, ROBIN E.; HERST, C. V.; KIES, MERRILL S.; MARDER, ROBERT J.; ROSEN, STEVEN T.

Archives of Pathology and Laboratory Medicine

November, 1987; 111: 1054-1056

LINE COUNT: 00122 WORD COUNT: 01684

2/3/18 (Item 18 from file: 442)

DIALOG(R) File 442: AMA Journals

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00037960

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Coexisting Hodgkin's Disease and Mycosis Fungoides; Immunohistochemical Proof of Its Existence (ORIGINAL ARTICLE)

SIMRELL, CHARLES R.; BOCCIA, RALPH V.; JAFFE, ELAINE S.

Archives of Pathology and Laboratory Medicine

November, 1986; 110: 1029-1034

LINE COUNT: 00223 WORD COUNT: 03078

2/3/19 (Item 19 from file: 442)

DIALOG(R) File 442: AMA Journals

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00029473

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Nephrotic Syndrome Associated With a Clonal T-Cell **Leukemia** of Large Granular Lymphocytes With Cytotoxic Function (CLINICAL OBSERVATION)

ORMAN, STEPHEN V.; SCHECHTER, GERALDINE P.; WHANG-PENG, JACQUELINE; GUCCION, JOHN; CHAN, CLARA; SCHULOF, RICHARD S.; SHALHOUB, ROBERT J.

Archives of Internal Medicine

September, 1986; 146: 1827-1829

LINE COUNT: 00157 WORD COUNT: 02170

2/3/20 (Item 20 from file: 442)

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00025675
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Sezary-like Syndrome in a 10-Year-Old Girl With Serologic Evidence of Human
T-Cell Lymphotropic Virus Type I Infection (OBSERVATIONS)
 IKAI, KOUICHI; UCHIYAMA, TAKASHI; MAEDA, MICHIYUKI; TAKIGAWA, MASAHIRO
 Archives of Dermatology
 October, 1987; 123: 1351-1355
 LINE COUNT: 00149
                           WORD COUNT: 02069
            (Item 21 from file: 442)
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DIALOG(R) File 442: AMA Journals
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Actinic Reticuloid: An Immunohistochemical Study (CORRESPONDENCE; Vignettes
)
 TAKIGAWA, MASAHIRO
 Archives of Dermatology
 March, 1987; 123: 296-297
 LINE COUNT: 00045
                    WORD COUNT: 00624
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           (Item 22 from file: 442)
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New Therapies for Cutaneous T-Cell Lymphoma (EDITORIALS)
  HEALD, PETER W.
  Archives of Dermatology
  February, 1987; 123: 189-191
  LINE COUNT: 00122
                           WORD COUNT: 01697
           (Item 23 from file: 442)
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Lymphomatoid Papulosis; Histologic and Immunohistochemical Studies
in a Patient With a Scaly Pigmented Eruption (STUDIES)
  TOKURA, YOSHIKI; TAKIGAWA, MASAHIRO; OKU, TOMOZO; YAMADA, MIZUHO
  Archives of Dermatology
  December, 1986; 122: 1400-1405
  LINE COUNT: 00215
                            WORD COUNT: 02972
            (Item 24 from file: 442)
DIALOG(R) File 442: AMA Journals
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Positron Emission Tomography in Oncology; Council on Scientific Affairs; Report of the Positron Emission Tomography Panel (Topics in Radiology/Council Report)

JACOBSON, HAROLD G.

JAMA, The Journal of the American Medical Association

April 8, 1988; 259: 2126-2131

LINE COUNT: 00216 WORD COUNT: 02988

2/3/25 (Item 25 from file: 442)

DIALOG(R) File 442:AMA Journals

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WYNGAARDEN, BY JAMES B.

JAMA, The Journal of the American Medical Association

October 2, 1987; 258: 1707

LINE COUNT: 00076 WORD COUNT: 01052

2/3/26 (Item 26 from file: 442)

DIALOG(R) File 442:AMA Journals

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Antitumor Strategies Based on Enhancing -- and Blocking -- Effects of Interleukin-2 (MEDICAL NEWS & PERSPECTIVES)

MERZ, BY BEVERLY

JAMA, The Journal of the American Medical Association

September 12, 1986; 256: 1241, 1244

LINE COUNT: 00114 WORD COUNT: 01577

2/3/27 (Item 1 from file: 444)

DIALOG(R) File 444: New England Journal of Med.

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00120088

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Prevention of Rejection in Cardiac Transplantation by Blockade of the Interleukin-2 Receptor with a Monoclonal Antibody (Original Articles)

Beniaminovitz, Ainat; Itescu, Silviu; Lietz, Katherine; Donovan, Mary; Burke, Elizabeth M.; Groff, Barbara D.; Edwards, Niloo; Mancini,

Donna M.

The New England Journal of Medicine

Mar 2, 2000; 342 (9),pp 613-619

LINE COUNT: 00418 WORD COUNT: 05780

2/3/28 (Item 2 from file: 444)

DIALOG(R) File 444: New England Journal of Med.

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00118604

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Bone Marrow Involvement in Acute **Leukemia** (Images in Clinical Medicine)

Junghans, R.P.
The New England Journal of Medicine
Nov 5, 1998; 339 (19),p 1375
LINE COUNT: 00074 WORD COUNT: 01028

2/3/29 (Item 3 from file: 444)
DIALOG(R)File 444:New England Journal of Med.
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00117715

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Interleukin-2-Receptor Blockade with Daclizumab to Prevent Acute Rejection in Renal Transplantation (Original Articles)

Vincenti, Flavio; Kirkman, Robert; Light, Susan; Bumgardner, Ginny; Pescovitz, Mark; Halloran, Philip; Neylan, John; Wilkinson, Alan; Ekberg, Henrik; Gaston, Robert; Backman, Lars; Burdick, James; for the Daclizumab Triple Therapy Study Group.
The New England Journal of Medicine
Jan 15, 1998; 338 (3),pp 161-165
LINE COUNT: 00291 WORD COUNT: 04020

2/3/30 (Item 4 from file: 444)
DIALOG(R)File 444:New England Journal of Med.
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00114651

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Treatment of Adult T-Cell Leukemia-Lymphoma with a Combination of Interferon Alfa and Zidovudine (Original Articles)

Gill, Parkash S.; Harrington, William, Jr.; Kaplan, Mark H.; Ribeiro, Raul C.; Bennett, John M.; Liebman, Howard A.; Bernstein-Singer, Marjorie; Espina, Byron M.; Cabral, Lisa; Allen, Steven; Kornblau, Steven; Pike, Malcolm C.; Levine, Alexandra M. The New England Journal of Medicine Jun 29, 1995; 332 (26),pp 1744-1748
LINE COUNT: 00357 WORD COUNT: 04935

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00113528

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Medical Progress: Renal Transplantation (Review Articles)

Suthanthiran, Manikkam; Strom, Terry B. The New England Journal of Medicine Aug 11, 1994; 331 (6),pp 365-376 LINE COUNT: 00629 WORD COUNT: 08682

2/3/32 (Item 6 from file: 444)
DIALOG(R)File 444:New England Journal of Med.
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00113464

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Current Concepts: Designer And Catalytic Antibodies (Review Article)

Mayforth, Ruth D.; Quintans, Jose. The New England Journal of Medicine Jul 19, 1990; 323 (3),pp 173-178

LINE COUNT: 00486 WORD COUNT: 06710

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Current Concepts: Designer And Catalytic Antibodies (Review Article)

Mayforth, Ruth D.; Quintans, Jose.
The New England Journal of Medicine
Jul 19, 1990; 323 (3),pp 173-178
LINE COUNT: 00486 WORD COUNT: 06710

2/3/34 (Item 8 from file: 444)
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Seminars in Medicine of the Beth Israel Hospital, Boston: Cytokine Receptors In Congenital Hematopoietic Disease (Review Article)

D'Andrea, Alan D.
The New England Journal of Medicine
Mar 24, 1994; 330 (12),pp 839-846
LINE COUNT: 00579 WORD COUNT: 08002

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00111325

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Seminars in Medicine of the Beth Israel Hospital, Boston: Pathogenesis Of Diseases Induced By Human Lymphotropic Virus Type I Infection (Review Articles)

Hollsberg, Per; Hafler, David A.
The New England Journal of Medicine
Apr 22, 1993; 328 (16),pp 1173-1182
LINE COUNT: 00561 WORD COUNT: 07742

2/3/36 (Item 10 from file: 444)
DIALOG(R)File 444:New England Journal of Med.
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Hodgkin's Disease, Lymphomatoid Papulosis, And Cutaneous T-Cell Lymphoma Derived From A Common T-Cell Clone (Original Articles)

Davis, Thomas H.; Morton, Cynthia C.; Miller-Cassman, Robert; Balk, Steven P.; Kadin, Marshall E.
The New England Journal of Medicine
Apr 23, 1992; 326 (17),pp 1115-1122
LINE COUNT: 00421 WORD COUNT: 05813

2/3/37 (Item 11 from file: 444)
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A Cluster Of Pneumocystis Carinii Pneumonia In Adults Without Predisposing Illnesses (Brief Report)

Jacobs, Jonathan L.; Libby, Daniel M.; Winters, Robert A.; Gelmont, David M.; Fried, Ethan D.; Hartman, Barry J.; Laurence, Jeffrey. The New England Journal of Medicine
Jan 24, 1991; 324 (4),pp 246-250
LINE COUNT: 00467 WORD COUNT: 06445

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Current Concepts: Designer And Catalytic Antibodies (Review Article)

Mayforth, Ruth D.; Quintas, Jose.
The New England Journal of Medicine
Jul 19, 1990; 323 (3),pp 173-178
LINE COUNT: 00487 WORD COUNT: 06722

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Severe Combined Immunodeficiency Due To A Specific Defect In The Production Of Interleukin-2 (Brief Reports)

Weinberg, Kenneth; Parkman, Robertson.
The New England Journal of Medicine
Jun 14, 1990; 322 (24),pp 1718-1723
LINE COUNT: 00466 WORD COUNT: 06442

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Weekly Clinicopathological Exercises: Case 36-1989: A 34-Year-Old Jamaican Man With Fever, Hepatic Failure, Diarrhea, And A Progressive Gait Disorder (Case Records of the Massachusetts General Hospital)

Groopman, Jerome E.; Ferry, Judith A.
The New England Journal of Medicine
Sep 7, 1989; 321 (10),pp 663-675
LINE COUNT: 01042 WORD COUNT: 14379

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00106327

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Pure Red-Cell Aplasia Of 10 Years' Duration Due To Persistent Parvovirus B19 Infection And Its Cure With Immunoglobulin Therapy (Medical Intelligence)

Kurtzman, Gary; Frickhofen, Norbert; Kimball, Janice; Jenkins, Douglas
W.; Nienhuis, Arthur W.; Young, Neal S.
The New England Journal of Medicine
Aug 24, 1989; 321 (8),pp 519-523
LINE COUNT: 00393 WORD COUNT: 05426

2/3/42 (Item 16 from file: 444)
DIALOG(R)File 444:New England Journal of Med.
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00104190

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Pathogenic Human Retroviruses (Editorial)

Broder, Samuel
The New England Journal of Medicine
January 28, 1988; 318 (4),pp 243-245
LINE COUNT: 00301 WORD COUNT: 04164

2/3/43 (Item 17 from file: 444) DIALOG(R)File 444:New England Journal of Med. (c) 2000 Mass. Med. Soc. All rts. reserv.

00102985

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Remissions In Hairy-Cell **Leukemia** With Pentostatin (2'-Deoxycoformycin) (Original Article)

Spiers, Alexander S.D., T.D., Ph.D.; Moore, Dirk, Ph.D., Cassileth, Peter A., Harrington, David P., Ph.D., Cummings, Frank J.; Neiman, Richard S., Bennett, John M.; O'Connell, Michael J.

The New England Journal of Medicine April 2, 1987; 316 (14),pp 825-830

LINE COUNT: 00439 WORD COUNT: 06069

2/3/44 (Item 18 from file: 444) DIALOG(R)File 444:New England Journal of Med. (c) 2000 Mass. Med. Soc. All rts. reserv.

00102420

Copyright 1986 by the Massachusetts Medical Society

Concomitant Infection with HTLV-I and HTLV-III In A Patient with T8

Lymphoproliferative Disease (Medical Intelligence) Harper, Mary E., Ph.D.; Kaplan, Mark H.; Marselle, Lisa M., B.A.; Pahwa, Savita G.; Chayt, Karen J.; Sarngadharan, M.G., Ph.D.; Wong-Staal, Flossie, Ph.D.; Gallo, Robert C. The New England Journal of Medicine October 23, 1986; 315 (17),pp 1073-1078 LINE COUNT: 00408 WORD COUNT: 05641 (Item 19 from file: 444) 2/3/45 DIALOG(R) File 444: New England Journal of Med. (c) 2000 Mass. Med. Soc. All rts. reserv. 00102091 Copyright 1986 by the Massachusetts Medical Society Second Isolate of HTLV-II Associated with Atypical Hairy-Cell Leukemia (Medical Intelligence) Ronseblatt, Joseph D.; Golde, David W.; Wachsman, William, Ph.D.; Giorgi, Janis V., Ph.D.; Jacobs, Andrew; Schmidt, Gerhard M.; Quan, Shirley; Gasson, Judith C., Ph.D.; Chen, Irvin S.Y., Ph.D. The New England Journal of Medicine August 7, 1986; 315 (6), pp 372-377 WORD COUNT: 07793 LINE COUNT: 00564 (Item 20 from file: 444) DIALOG(R) File 444: New England Journal of Med. (c) 2000 Mass. Med. Soc. All rts. reserv. 00101373 Copyright 1986 by the Massachusetts Medical Society Isolate

of HTLV-II Associated with Atypical Hairy-Cell Leukemia (Medical Intelligence)

Ronseblatt, Joseph D.; Golde, David W.; Wachsman, William, Ph.D.; Giorgi, Janis V., Ph.D.; Jacobs, Andrew; Schmidt, Gerhard M.; Quan, Shirley; Gasson, Judith C., Ph.D.; Chen, Irvin S.Y., Ph.D. The New England Journal of Medicine August 7, 1986; 315 (6),pp 372-377 LINE COUNT: 00564 WORD COUNT: 07793

(Item 21 from file: 444) DIALOG(R) File 444: New England Journal of Med. (c) 2000 Mass. Med. Soc. All rts. reserv.

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T-Cell Receptor Gene Rearrangements as Clinical Markers of Human T-Cell Lymphomas (Original Article)

Bertness, Virginia, B.S.; Kirsch, Ilan; Hollis, Gregory, Ph.D.; Johnson, Bruce; Bunn, Paul A., Jr. The New England Journal of Medicine August 29, 1985; 313 (9),pp 534-538 LINE COUNT: 00579 WORD COUNT: 07994

(Item 22 from file: 444) DIALOG(R) File 444: New England Journal of Med. (c) 2000 Mass. Med. Soc. All rts. reserv.

00100384 Copyright 1985 by the Massachusetts Medical Society

Subsets of Patients with Aplastic Anemia Identified by Flow Microfluorometry (Original Article)

Torok-Storb, Beverly, Ph.D.; Doney, Kristine; Sale, George; Thomas, E. Donnall; Storb, Rainer
The New England Journal of Medicine
April 18, 1985; 312 (16),pp 1015-1022
LINE COUNT: 00484 WORD COUNT: 06684

2/3/49 (Item 1 from file: 457)
DIALOG(R)File 457:The Lancet
(c) 2000 The Lancet, Ltd. All rts. reserv.

00118363 (USE FORMAT 7 OR 9 FOR FULLTEXT)
TITLE: Human T-lymphotropic virus type I infection
Manns, Angela; Hisada, Michie; La Grenade, Lois
Viral Epidemiology Branch, Division of Cancer Epidemiology and Genetics,
National Cancer Institute, Bethesda, MD, USA; Department of Medicine,
University of the West Indies, Kingston, Jamaica
The Lancet, v353, n9168, pp 1951-1958
June 5, 1999

DOCUMENT TYPE: Journal; Seminar LANGUAGE: English RECORD TYPE: Fulltext
WORD COUNT: 5001

2/3/50 (Item 2 from file: 457)
DIALOG(R)File 457:The Lancet
(c) 2000 The Lancet, Ltd. All rts. reserv.

00117541 (USE FORMAT 7 OR 9 FOR FULLTEXT)
TITLE: Immunosuppressive strategies in transplantation
Denton, Mark D; Magee, Colm C; Sayegh, Mohamed H
Laboratory of Immunogenetics and Transplantation, Renal Division,
Department of Medicine, Brigham and Women's Hospital, Harvard Medical
School, Boston, MA 02115, USA
The Lancet, v353, n9158, pp 1083-1091
March 27, 1999

DOCUMENT TYPE: Journal; Review LANGUAGE: English RECORD TYPE: Fulltext WORD COUNT: 5609

2/3/51 (Item 3 from file: 457)
DIALOG(R)File 457:The Lancet
(c) 2000 The Lancet, Ltd. All rts. reserv.

00104852 (USE FORMAT 7 OR 9 FOR FULLTEXT)
TITLE: Cutaneous T-cell lymphoma (mycosis fungoides)
Lorincz, Allan L
Section of Dermatology, University of Chicago, 5841 South Maryland, MC 5067, Chicago, IL 60637, USA
The Lancet, v347, n9005, pp 871-876
March 30, 1996

DOCUMENT TYPE: Journal; Skin cancer quartet LANGUAGE: English RECORD TYPE: Fulltext

WORD COUNT: 3304

2/3/52 (Item 4 from file: 457) DIALOG(R) File 457: The Lancet (c) 2000 The Lancet, Ltd. All rts. reserv. 00101288 (USE FORMAT 7 OR 9 FOR FULLTEXT) TITLE: Impairment of leukaemia-free survival by addition of interleukin-2receptor antibody to standard graft-versus-host prophylaxis Blaise, Didier; Olive, Daniel; Michallet, Mauricette; Marit, Gerald; Leblond, Veronique; Maraninchi, Dominique Institut Paoli Calmettes and Inserm U119, Marseille; Hopital Edouard Herriot, Lyon; Hopital Haut Leveque, Bordeaux, Hopital Pitie-Salpetriere, Paris, France. The Lancet, v345, n8958, pp 1144-1146 May 6, 1995 DOCUMENT TYPE: Journal; Articles LANGUAGE: English RECORD TYPE: Fulltext WORD COUNT: 1856 2/3/53 (Item 5 from file: 457) DIALOG(R) File 457: The Lancet (c) 2000 The Lancet, Ltd. All rts. reserv. 00088561 (USE FORMAT 7 OR 9 FOR FULLTEXT) TITLE: Short Reports: Therapeutic effects of genetically engineered toxin (DAB sub 486 IL-2) in patient with chronic lymphocytic leukaemia LeMaistre, Charles F|Rosenblum, Michael G|Reuben, James M|Parkinson, David R|Meneghetti, Carole M|Parker, Karen|Shaw, Jill P|Deisseroth, Albert B|Woodworth, Thasia Department of Medicine/Hematology, University of Texas Health Science Center, San Antonio. Department of Hematology, Department of Clinical Immunology an Biological Therapy and Department of Laboratory Medicine, University of Texas MD Anderson Cancer Centre, Houston, Texas. National Cancer Institute, Bethesda, Maryland. Seragen, Inc., Hopkinton, Massachusetts, USA. Correspondence: Dr C. F. LeMaistre, University of Texas Health Science Center, Department of Medicine/Hematology, 7703 Floyd Curl Drive, San Antonio, Texas 78284-7880, USA. The Lancet, v337, n8750, pp 1124-1125 1991 May 11 DOCUMENT TYPE: Journal; Report (REP) LANGUAGE: English RECORD TYPE: Fulltext WORD COUNT: 993 (Item 6 from file: 457) DIALOG(R) File 457: The Lancet (c) 2000 The Lancet, Ltd. All rts. reserv. 00082387 (USE FORMAT 7 OR 9 FOR FULLTEXT) TITLE: Original Article: Effects of intradermal gamma-interferon in cutaneous leishmaniasis Harms, G|Zwingenberger, K|Chehade, A K|Talhari, S|Racz, P|Mouakeh, A|Douba, M|Nakel, L|Naiff, R D|Kremsner, P G|Feldmeier, H|Bienzle, U Landesinstitut fur Tropenmedizin, Berlin, Federal Republic of Germany. Departement de Dermato-Venereologie, Faculte de Medecine, Universite d'Alep, Syria. Centro de Dermatologia Tropical e Venereologia "Alfredo da Matta", Manaus, Brazil. Bernhard-Nocht-Institut fur Tropenmedizin, Abteilung fur Pathologie, Hamburg, Federal Republic of Germany. Instituto Nacional de Pesquisa da Amazonia, Departamento de Leishmaniose, Manaus, Brazil. Correspondence: G. H., Landesinstitut fur Tropenmedizin,

Koenigin-Elisabeth-Strasse 32, 1 Berlin 19, FRG.

The Lancet, v333, n8650, pp 1287-1292

1989 Jun 10

DOCUMENT TYPE: Journal; Article (ART) LANGUAGE: English RECORD TYPE:

Fulltext

WORD COUNT: 3343

2/3/55 (Item 7 from file: 457)

DIALOG(R) File 457: The Lancet

(c) 2000 The Lancet, Ltd. All rts. reserv.

00075191 (USE FORMAT 7 OR 9 FOR FULLTEXT)

TITLE: Original Articles: Serum reactivity to human T-cell leukaemia/ lymphoma virus type I proteins in patients with large granular lymphocytic leukaemia

Starkebaum, Gordon|Loughran, Thomas P Jr|Kalyanaraman, V S|Kadin, Marshall E|Kidd, Pamela G|Singer, Jack W|Ruscetti, Francis W

Department of Medicine, Veterans Administration Medical Center, Seattle, Washington. Fred Hutchinson Cancer Research Center, Seattle. Centers for Disease Control, Atlanta, Georgia. Department of Pathology and Charles A. Dana Research Laboratory, Beth Israel Hospital and Harvard Medical School, Boston, Massachusetts. Department of Medicine and Laboratory Medicine, University of Washington, Seattle. National Cancer Institute, Frederick Cancer Research Facility, Frederick, Maryland, USA. Correspondence: G. S., Medical Service (111), VA Medical Center, 1660 S. Columbian Way, Seattle, Washington 98108, USA.

The Lancet, v329, n8533, pp 596-598 1987 Mar 14

DOCUMENT TYPE: Journal; Article (ART) LANGUAGE: English RECORD TYPE:

Fulltext

WORD COUNT: 1984 ? s s1 and py=1993

78 S1 9725 PY=1993

S3 2 S1 AND PY=1993

? t s3/3/all

3/3/1 (Item 1 from file: 442)
DIALOG(R)File 442:AMA Journals
(c)2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

00087400

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(ARTICLE)

GOLDMAN, BARRY D.; OH, SE-KYUNG; DAVIS, BRET E.; KADIN, MARSHALL E.; POIESZ, BERNARD J.; KOH, HOWARD K.

Archives of Dermatology

Sep, 1993; OBSERVATION: p1166

LINE COUNT: 00294

3/3/2 (Item 1 from file: 444)
DIALOG(R)File 444:New England Journal of Med.
(c) 2000 Mass. Med. Soc. All rts. reserv.

00111325

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Seminars in Medicine of the Beth Israel Hospital, Boston: Pathogenesis Of Diseases Induced By Human Lymphotropic Virus Type I Infection (Review Articles)

Hollsberg, Per; Hafler, David A. The New England Journal of Medicine Apr 22, **1993**; 328 (16),pp 1173-1182

LINE COUNT: 00561 WORD COUNT: 07742 ? s s1 and py=1994

78 S1 9636 PY=1994 4 S1 AND PY=1994 S4

? t s4/3/all

4/3/1 (Item 1 from file: 442) DIALOG(R) File 442:AMA Journals (c) 2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

00090827

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Recurrence of Pemphigus Vulgaris Associated With Interleukin 2 Therapy (ARTICLE)

PRUSSICK, RONALD; PLOTT, R. TODD; STANLEY, JOHN R.

Archives of Dermatology

July, 1994; Observation: de 890

LINE COUNT: 00266

4/3/2 (Item 2 from file: 442) DIALOG(R) File 442:AMA Journals (c) 2000 Amer Med Assn -FARS/DARS apply. All rts. reserv.

00090215

COPYRIGHT American Medical Association 1994

Primary Gastric T-Cell Lymphoma Morphological and Immunohistochemical Studies of Two Cases (ARTICLE)

Archives of Pathology and Laboratory Medicine May, 1994; Original Article: p547

LINE COUNT: 00257

(Item 1 from file: 444) DIALOG(R) File 444: New England Journal of Med. (c) 2000 Mass. Med. Soc. All rts. reserv.

00113528

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Medical Progress: Renal Transplantation (Review Articles)

Suthanthiran, Manikkam; Strom, Terry B. The New England Journal of Medicine Aug 11, **1994**; 331 (6),pp 365-376

LINE COUNT: 00629 WORD COUNT: 08682

(Item 2 from file: 444) DIALOG(R) File 444: New England Journal of Med. (c) 2000 Mass. Med. Soc. All rts. reserv.

00112704

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Seminars in Medicine of the Beth Israel Hospital, Boston: Cytokine Receptors In Congenital Hematopoietic Disease (Review Article)

D'Andrea, Alan D.

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LINE COUNT: 00579
                            WORD COUNT: 08002
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       (c) format only 2000 Dialog Corporation
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The New England Journal of Medicine Mar 24, 1994; 330 (12),pp 839-846

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4 AU=WALDMAN TA
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      S1
             254 E1, E19
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72 S1 AND TAC
? s s2 and py=1993
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80574 PY=1993
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? s s2 and py=1994
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83740 PY=1994
              4 S2 AND PY=1994
? t s3/3/a11
 3/3/1
DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
01034122
           94003208
                      MEDL/94003208
  The interleukin-2 receptor: a target for monoclonal antibody treatment of
human T-cell lymphotrophic virus I-induced adult T-cell leukemia.
  Waldmann TA; White JD; Goldman CK; Top L; Grant A; Bamford R;
Roessler E; Horak ID; Zaknoen S; Kasten-Sportes C; et al
  Metabolism Branch and Laboratory of Pathology, National Cancer Institute,
National Institutes of Health, Bethesda, MD 20892.
  Blood; 82(6):1701-12 1993 ISSN 0006-4971
                                              Journal Code: A8G
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE
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DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
           93368196
                     MEDL/93368196
  1992 Stohlman Memorial Lecture: targeting the IL-2 receptor.
  Waldmann TA
  Metabolism Branch, National Cancer Institute, National Institutes of
Health, Bethesda, MD 20892.
  Leukemia; 7 Suppl 2:S151-6 1993 ISSN 0887-6924 Journal Code: LEU
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE
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DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
         93329041 MEDL/93329041
  Humanized Mik beta 1, a humanized antibody to the IL-2 receptor
beta-chain that acts synergistically with humanized anti-TAC.
Hakimi J; Ha VC; Lin P; Campbell E; Gately MK; Tsudo M; Payne PW; Waldmann TA; Grant AJ; Tsien WH; et al
  Roche Research Center, Hoffmann-LaRoche, Inc., Nutley, NJ 07110.
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J Immunol; 151(2):1075-85 1993 ISSN 0022-1767 Journal Code: IFB

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

3/3/4 DIALOG(R) File 159: Cancerlit (c) format only 2000 Dialog Corporation. All rts. reserv. 93218273 MEDL/93218273 Cytotoxic activities of recombinant immunotoxins composed of Pseudomonas toxin or diphtheria toxin toward lymphocytes from patients with adult T-cell leukemia. Kreitman RJ; Chaudhary VK; Waldmann TA; Hanchard B; Cranston B; FitzGerald DJ; Pastan I Laboratory of Molecular Biology, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892. Leukemia; 7(4):553-62 1993 ISSN 0887-6924 Journal Code: LEU Languages: ENGLISH Document Type: JOURNAL ARTICLE 3/3/5 DIALOG(R) File 159: Cancerlit (c) format only 2000 Dialog Corporation. All rts. reserv. 95606931 ICDB/95606931 Lymphokine receptors: a target for immunotherapy of lymphomas (Meeting abstract). Waldmann TA NCI, Bethesda, MD 20892 Non-serial; Fifth International Conference on Malignant Lymphoma, June 9-12, 1993, Lugano, Switzerland, p. 46, 1993.: 1993 Languages: ENGLISH Document Type: MEETING ABSTRACTS 3/3/6 DIALOG(R) File 159: Cancerlit (c) format only 2000 Dialog Corporation. All rts. reserv. 00985141 93693792 ICDB/93693792 Towards optimal anti-Tac antibody dosing in the presence of antiqenemia in adult T-cell leukemia-lymphoma (Meeting abstract). RP; Goldman CK; Carrasquillo J; Reynolds J; Nelson DL; Waldmann TA New England Deaconess Hosp., Boston, MA 02215 Proc Annu Meet Am Assoc Cancer Res; 34:A2849 1993 ISSN 0197-016X Languages: ENGLISH Document Type: MEETING ABSTRACTS ? t s4/3/all 4/3/1 DIALOG(R) File 159: Cancerlit (c) format only 2000 Dialog Corporation. All rts. reserv. 01102643 94325579 MEDL/94325579

Treatment of acute graft-versus-host disease with humanized anti-Tac: an antibody that binds to the interleukin-2 receptor.

Anasetti C; Hansen JA; Waldmann TA; Appelbaum FR; Davis J; Deeg HJ; Doney K; Martin PJ; Nash R; Storb R; et al

Clinical Research Division, Fred Hutchinson Cancer Research Center 98104. Blood; 84(4):1320-7 1994 ISSN 0006-4971 Journal Code: A8G

Contract/Grant No.: CA 18029, SS, CA, SS, NCI; CA 18221, SS, CA, SS, NCI; HL36444, SS, HL, SS, NHLBI; +

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

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DIALOG(R)File 159:Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
01092742
           94266289 MEDL/94266289
   Anti-IL-2 receptor monoclonal antibody (anti-Tac) treatment of
T-cell lymphoma.
  Waldmann TA
  Metabolism Branch, National Cancer Institute, Bethesda, Maryland.
  Important Adv Oncol; :131-41 1994 ISSN 0883-5896 Journal Code: GG9
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, ACADEMIC
 4/3/3
DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
           94226936
                    MEDL/94226936
  Lymphokine receptors: a target for immunotherapy of lymphomas.
  Waldmann TA
  Metabolism Branch, National Cancer Institute, National Institutes of
Health, Bethesda, Maryland.
  Ann Oncol; 5 Suppl 1:13-7 1994 ISSN 0923-7534 Journal Code: AYF
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL
 4/3/4
DIALOG(R)File 159:Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
01083831
           94211860
                     MEDL/94211860
  Cooperative interactions between the interleukin 2 receptor alpha and
beta chains alter the interleukin 2-binding affinity of the receptor
subunits.
  Roessler E; Grant A; Ju G; Tsudo M; Sugamura K; Waldmann TA
  Metabolism Branch, National Cancer Institute, National Institutes of
Health, Bethesda, MD 20892.
  Proc Natl Acad Sci U S A; 91(8):3344-7 1994 ISSN 0027-8424
Journal Code: PV3
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE
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S1
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     S5
              69 RD S2 (unique items)
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 5/3/1
DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
01559550
          20021681
                    MEDL/20021681
  Responses in refractory hairy cell leukemia to a recombinant immunotoxin.
  Kreitman RJ; Wilson WH; Robbins D; Margulies I; Stetler-Stevenson M;
Waldmann TA; Pastan I
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Laboratory of Molecular Biology, the Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, MD.

Blood; 94(10):3340-8 1999 ISSN 0006-4971 Journal Code: A8G

Languages: ENGLISH

Document Type: CLINICAL TRIAL; CLINICAL TRIAL, PHASE I; JOURNAL ARTICLE

5/3/2

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01516566 99251477 MEDL/99251477

Favorable effects of glycolate conjugation on the biodistribution of humanized antiTac Fab fragment.

Kobayashi H; Kim IS; Drumm D; Kim MK; Paik DS; Le N; Waldmann TA; Carrasquillo JA; Paik CH

Department of Nuclear Medicine, Warren G. Magnuson Clinical Center, National Institutes of Health, Bethesda, Maryland 20892-1180, USA.

J Nucl Med; 40(5):837-45 1999 ISSN 0161-5505 Journal Code: JEC

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/3

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01479275 99066764 MEDL/99066764

Reduction in HTLV-I proviral load and spontaneous lymphoproliferation in HTLV-I-associated myelopathy/tropical spastic paraparesis patients treated with humanized anti-Tac.

Lehky TJ; Levin MC; Kubota R; Bamford RN; Flerlage AN; Soldan SS; Leist TP; Xavier A; White JD; Brown M; Fleisher TA; Top LE; Light S; McFarland HF; Waldmann TA; Jacobson S

Neuroimmunology Branch, National Institute of Neurological Disorders and Stroke, National Institutes of Health, Bethesda, MD, USA.

Ann Neurol; 44(6):942-7 1998 ISSN 0364-5134 Journal Code: 6AE

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/4

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01441042 98301301 MEDL/98301301

Examination of a role for idiotypy in the disease remission of a long-term survivor of adult T cell leukemia treated with anti-Tac antibody.

Kingsbury GA; Waldmann TA; Junghans RP

Biotherapeutics Development Lab, Harvard Medical School, Beth Israel Deaconess Medical Center, Boston, MA 02115, USA.

Leukemia; 12(6):982-91 1998 ISSN 0887-6924 Journal Code: LEU

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/5

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01436925 98280344 MEDL/98280344

Epitope blocking: positive and negative effects on the biodistribution of 125I-labeled anti-Tac disulfide-stabilized Fv fragment of two antibodies against different epitopes of the circulating antigen.

Kobayashi H; Sun BF; Han ES; Kim MK; Le N; Wang QC; Nelson DL; Pastan I;

Waldmann TA; Paik CH; Carrasquillo JA

Department of Nuclear Medicine, Warren G. Magnuson Clinical Center, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892-1180, USA.

Jpn J Cancer Res; 89(4):436-44 1998 ISSN 0910-5050 Journal Code: HBA

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/6

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01414399 98132664 MEDL/98132664

Impact of antigenemia on the bioactivity of infused anti-Tac antibody: implications for dose selection in antibody immunotherapies.

Junghans RP; Carrasquillo JA; Waldmann TA

Biotherapeutics Development Lab, Harvard Medical School, Division of Hematology-Oncology, Beth Israel Deaconess Medical Center, Boston, MA 02115, USA. junghans@warren.med.harvard.edu Proc Natl Acad Sci U S A; 95(4):1752-7 1998 ISSN 0027-8424

Journal Code: PV3 Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/7

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01353147 97301590 MEDL/97301590

Improved biodistribution of 125I-labeled disulfide-stabilized Fv fragment by blocking its binding to the alpha subunit of the interleukin 2 receptor in the circulation with preinjected humanized anti-Tac IgG.

Kobayashi H; Yoo TM; Drumm D; Kim MK; Sun BF; Le N; Webber KO; Pastan I; Waldmann TA; Paik CH; Carrasquillo JA

Department of Nuclear Medicine, National Cancer Institute, NIH, Bethesda, Maryland 20892-1180, USA.

Cancer Res; 57(10):1955-61 1997 ISSN 0008-5472 Journal Code: CNF

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/8

DIALOG(R) File 159: Cancerlit

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01326403 98641089 ICDB/98641089

Phase I clinical trial with recombinant immunotoxin anti-Tac (Fv)-PE38 (LMB-2) in patients with hematologic malignancies (Meeting

Kreitman RJ; White JD; Pearson D; Top LE; Waldmann TA; Pastan I National Cancer Institute, National Institutes of Health, Bethesda, MD 20892-4255

Proc Annu Meet Am Assoc Cancer Res; 38:A4089 1997 ISSN 0197-016X Languages: ENGLISH

Document Type: MEETING ABSTRACTS; CLINICAL TRIAL; CLINICAL TRIAL, PHASE I

5/3/9

DIALOG(R) File 159: Cancerlit

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01281441 96390748 MEDL/96390748

The promiscuous IL-2/IL-15 receptor: a target for immunotherapy of

HTLV-I-associated disorders.

Waldmann TA

Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, Maryland 20892, USA.

J Acquir Immune Defic Syndr Hum Retrovirol; 13 Suppl 1:S179-85 1996 ISSN 1077-9450 Journal Code: B7J

Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL

5/3/10

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01261511 96261660 MEDL/96261660

Metabolism of **Tac** (IL2Ralpha): physiology of cell surface shedding and renal catabolism, and suppression of catabolism by antibody binding. Junghans RP; **Waldmann TA**

Division of Hematology-Oncology, Harvard Medical School, New England Deaconess Hospital, Boston, MA 02215, USA.

J Exp Med; 183(4):1587-602 1996 ISSN 0022-1007 Journal Code: I2V

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/11

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01213309 96082159 MEDL/96082159

Radioimmunotherapy of interleukin-2R alpha-expressing adult T-cell leukemia with Yttrium-90-labeled anti-Tac [see comments]

Waldmann TA; White JD; Carrasquillo JA; Reynolds JC; Paik CH; Gansow OA; Brechbiel MW; Jaffe ES; Fleisher TA; Goldman CK; et al

Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892, USA.

Blood; 86(11):4063-75 1995 ISSN 0006-4971 Journal Code: A8G

Comment in SS Blood SS 1996 Jun 15;87(12):5379-80

Languages: ENGLISH

Document Type: CLINICAL TRIAL; CLINICAL TRIAL, PHASE I; CLINICAL TRIAL, PHASE II; JOURNAL ARTICLE

5/3/12

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01170001 95241460 MEDL/95241460

Phenotypic knockout of the high-affinity human interleukin 2 receptor by intracellular single-chain antibodies against the alpha subunit of the receptor.

Richardson JH; Sodroski JG; Waldmann TA; Marasco WA

Department of Pathology, Dana-Farber Cancer Institute, Harvard Medical School, Boston, MA 02115, USA.

Proc Natl Acad Sci U S A; 92(8):3137-41 1995 ISSN 0027-8424

Journal Code: PV3

Contract/Grant No.: P30 AI28691, SS, AI, SS, NIAID; P30 CA06516, SS, CA, SS, NCI; AI28785, SS, AI, SS, NIAID; +

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/13

DIALOG(R) File 159: Cancerlit

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01156141 97609760 ICDB/97609760

The promiscuous IL-2 receptor: a target for immunotherapy of HTLV-I-associated diseases (Meeting abstract).

Waldmann TA

Metabolism Branch, NCI, NIH Bethesda, MD 20892 Int J Oncol; 7(Suppl):980 1995 ISSN 1019-6439

Languages: ENGLISH

Document Type: MEETING ABSTRACTS

5/3/14

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01102643 94325579 MEDL/94325579

Treatment of acute graft-versus-host disease with humanized anti-Tac: an antibody that binds to the interleukin-2 receptor.

Anasetti C; Hansen JA; Waldmann TA; Appelbaum FR; Davis J; Deeg HJ;

Doney K; Martin PJ; Nash R; Storb R; et al

Clinical Research Division, Fred Hutchinson Cancer Research Center 98104.

Blood; 84(4):1320-7 1994 ISSN 0006-4971 Journal Code: A8G

Contract/Grant No.: CA 18029, SS, CA, SS, NCI; CA 18221, SS, CA, SS, NCI; HL36444, SS, HL, SS, NHLBI; +

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/15

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01092742 94266289 MEDL/94266289

Anti-IL-2 receptor monoclonal antibody (anti- ${\bf Tac}$) treatment of T-cell lymphoma.

Waldmann TA

Metabolism Branch, National Cancer Institute, Bethesda, Maryland. Important Adv Oncol; :131-41 1994 ISSN 0883-5896 Journal Code: GG9 Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, ACADEMIC

5/3/16

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01086036 94226936 MEDL/94226936

Lymphokine receptors: a target for immunotherapy of lymphomas.

Waldmann TA

Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, Maryland.

Ann Oncol; 5 Suppl 1:13-7 1994 ISSN 0923-7534 Journal Code: AYF

Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL

5/3/17

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01083831 94211860 MEDL/94211860

Cooperative interactions between the interleukin 2 receptor alpha and beta chains alter the interleukin 2-binding affinity of the receptor subunits.

Roessler E; Grant A; Ju G; Tsudo M; Sugamura K; Waldmann TA Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892.

Proc Natl Acad Sci U S A; 91(8):3344-7 1994 ISSN 0027-8424

Journal Code: PV3
Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/18

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01034122 94003208 MEDL/94003208

The interleukin-2 receptor: a target for monoclonal antibody treatment of human T-cell lymphotrophic virus I-induced adult T-cell leukemia.

Waldmann TA; White JD; Goldman CK; Top L; Grant A; Bamford R; Roessler E; Horak ID; Zaknoen S; Kasten-Sportes C; et al

Metabolism Branch and Laboratory of Pathology, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892.

Blood; 82(6):1701-12 1993 ISSN 0006-4971 Journal Code: A8G

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/19

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01029144 93368196 MEDL/93368196

1992 Stohlman Memorial Lecture: targeting the IL-2 receptor.

Waldmann TA

Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892.

Leukemia; 7 Suppl 2:S151-6 1993 ISSN 0887-6924 Journal Code: LEU

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/20

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01022892 93329041 MEDL/93329041

Humanized Mik beta 1, a humanized antibody to the IL-2 receptor beta-chain that acts synergistically with humanized anti-TAC.

Hakimi J; Ha VC; Lin P; Campbell E; Gately MK; Tsudo M; Payne PW; Waldmann TA; Grant AJ; Tsien WH; et al

Roche Research Center, Hoffmann-LaRoche, Inc., Nutley, NJ 07110. J Immunol; 151(2):1075-85 1993 ISSN 0022-1767 Journal Code: IFB

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/21

DIALOG(R) File 159: Cancerlit

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01004177 93218273 MEDL/93218273

Cytotoxic activities of recombinant immunotoxins composed of Pseudomonas toxin or diphtheria toxin toward lymphocytes from patients with adult T-cell leukemia.

Kreitman RJ; Chaudhary VK; Waldmann TA; Hanchard B; Cranston B;
FitzGerald DJ; Pastan I

Laboratory of Molecular Biology, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892.

Leukemia; 7(4):553-62 1993 ISSN 0887-6924 Journal Code: LEU

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/22

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00985141 93693792 ICDB/93693792

Towards optimal anti-Tac antibody dosing in the presence of antigenemia in adult T-cell leukemia-lymphoma (Meeting abstract).

Junghans RP; Goldman CK; Carrasquillo J; Reynolds J; Nelson DL; Waldmann TA

New England Deaconess Hosp., Boston, MA 02215

Proc Annu Meet Am Assoc Cancer Res; 34:A2849 1993 ISSN 0197-016X

Languages: ENGLISH

Document Type: MEETING ABSTRACTS

5/3/23

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00970994 93102188 MEDL/93102188

Prolongation of graft survival in primate allograft transplantation by yttrium-90-labeled anti-**Tac** in conjunction with granulocyte colony-stimulating factor.

Parenteau GL; Dirbas FM; Garmestani K; Brechbiel MW; Bukowski MA; Goldman CK; Clark R; Gansow OA; Waldmann TA

Surgery Branch, National Heart, Lung and Blood Institute, National Institutes of Health, Bethesda, Maryland 20892.

Transplantation; 54(6):963-8 1992 ISSN 0041-1337 Journal Code: WEJ

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/24

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00961028 93043351 MEDL/93043351

Recombinant toxins containing the variable domains of the anti-Tac monoclonal antibody to the interleukin-2 receptor kill malignant cells from patients with chronic lymphocytic leukemia.

Kreitman RJ; Chaudhary VK; Kozak RW; FitzGerald DJ; Waldman TA; Pastan I

Laboratory of Molecular Biology, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892.

Blood; 80(9):2344-52 1992 ISSN 0006-4971 Journal Code: A8G

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/25

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00889798 92035744 MEDL/92035744

Lymphokine receptor-directed therapy: a model for immune intervention in leukemia, autoimmunity, and immunodeficiency.

Waldmann TA

Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, Maryland 20892.

Clin Immunol Immunopathol; 61(2 Pt 2):S37-46 1991 ISSN 0090-1229 Journal Code: DEA

Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL

5/3/26

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00847429 92678733 ICDB/92678733

ADULT T-CELL LEUKEMIA: PROSPECTS FOR IMMUNOTHERAPY

Waldmann TA

Metabolism Branch, NCI, Bethesda, MD 20892

Non-serial; The Human Retroviruses. Gallo RC and Jay G, eds. San Diego, Academic Press, p. 319-32, 1991.: 1991

Languages: ENGLISH

Document Type: MONOGRAPH; REVIEW

5/3/27

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00792675 90205200 MEDL/90205200

Human B lymphocytes express the p75 component of the interleukin 2 receptor.

Begley CG; Burton JD; Tsudo M; Brownstein BH; Ambrus JL Jr; Waldmann TA

Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892.

Leuk Res; 14(3):263-71 1990 ISSN 0145-2126 Journal Code: K9M

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/28

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00787046 90150108 MEDL/90150108

Anti-Tac-H, a humanized antibody to the interleukin 2 receptor with new features for immunotherapy in malignant and immune disorders. Junghans RP; Waldmann TA; Landolfi NF; Avdalovic NM; Schneider WP;

Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, Maryland 20892.

Cancer Res; 50(5):1495-502 1990 ISSN 0008-5472 Journal Code: CNF

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/29

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DIALOG(R)File 159:Cancerlit

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00778698 90367069 MEDL/90367069

 $\ensuremath{\text{IL-2}}$ receptor expression in the haematologic malignancies: a target for immunotherapy.

Waldmann TA

Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, Maryland.

Cancer Surv; 8(4):891-903 1989 ISSN 0261-2429 Journal Code: CNG

Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL

5/3/30

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00762583 90053574 MEDL/90053574

The multichain interleukin-2 receptor: a target for immunotherapy of patients receiving allografts.

Waldmann TA; Goldman CK

Metabolism Branch National Cancer Institute, Bethesda, MD 20892.

Am J Kidney Dis; 14(5 Suppl 2):45-53 1989 ISSN 0272-6386

Journal Code: 3H5

Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL

5/3/31

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00758904 90028731 MEDL/90028731

Comparison of anti-Tac and anti-transferrin receptor-conjugated liposomes for specific drug delivery to adult T-cell leukemia.

Hege KM; Daleke DL; Waldmann TA; Matthay KK

Department of Pediatrics, University of California, San Francisco 94143.

Blood; 74(6):2043-52 1989 ISSN 0006-4971 Journal Code: A8G

Contract/Grant No.: CA39448, SS, CA, SS, NCI

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/32

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00746870 89337215 MEDL/89337215

The interaction of interleukin 2 with its receptor in the generation of suppressor T cells in antigen-specific and antigen-nonspecific systems in vitro.

Oh-Ishi T; Goldman CK; Misiti J; Waldmann TA

Metabolism Branch, National Cancer Institute, Bethesda, Maryland 20892.

Clin Immunol Immunopathol; 52(3):447-59 1989 ISSN 0090-1229

Journal Code: DEA Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/33

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00738209 89279941 MEDL/89279941

Multichain interleukin-2 receptor: a target for immunotherapy in lymphoma.

Waldmann TA

Metabolism Branch, National Cancer Institute, Bethesda, MD 20892.

J Natl Cancer Inst; 81(12):914-23 1989 ISSN 0027-8874 Journal Code: J9J

Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, ACADEMIC

5/3/34

DIALOG(R) File 159: Cancerlit

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00732036 89230284 MEDL/89230284

Nature of the bifunctional chelating agent used for radioimmunotherapy with yttrium-90 monoclonal antibodies: critical factors in determining in vivo survival and organ toxicity.

Kozak RW; Raubitschek A; Mirzadeh S; Brechbiel MW; Junghaus R; Gansow OA;

Waldmann TA

Division of Cytokine Biology, Center for Biologics Evaluation and Research, FDA, Bethesda, Maryland.

Cancer Res; 49(10):2639-44 1989 ISSN 0008-5472 Journal Code: CNF

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/35

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00731058 89223365 MEDL/89223365

Early experience with anti-Tac in clinical renal transplantation.

Kirkman RL; Shapiro ME; Carpenter CB; Milford EL; Ramos EL; Tilney NL; Waldmann TA; Zimmerman CE; Strom TB

Department of Surgery, Brigham and Women's Hospital, Boston, Massachusetts 02115.

Transplant Proc; 21(1 Pt 2):1766-8 1989 ISSN 0041-1345 Journal Code: WE9

Languages: ENGLISH

Document Type: CLINICAL TRIAL; JOURNAL ARTICLE

5/3/36

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00729030 89203989 MEDL/89203989

Anti-TAC MOAB prolongs renal allografts in cynomolgus monkeys.

Reed MH; Shapiro ME; Strom TB; Carpenter CB; Letvin NL; Reimann K; Weinberg DS; Waldmann TA; Kirkman RL

Brigham and Women's Hospital, Harvard Medical School, Boston, Massachusetts 02115.

Transplant Proc; 21(1 Pt 1):1028-30 1989 ISSN 0041-1345 Journal Code:

Contract/Grant No.: N01-AI-52587, SS, AI, SS, NIAID; P01-AI-19414, SS, AI, SS, NIAID; RR-01032, SS, RR, SS, NCRR

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/37

DIALOG(R) File 159: Cancerlit

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00694691 89027118 MEDL/89027118

Therapy of patients with human T-cell lymphotrophic virus I-induced adult T-cell leukemia with anti-Tac, a monoclonal antibody to the receptor for interleukin-2.

Waldmann TA; Goldman CK; Bongiovanni KF; Sharrow SO; Davey MP; Cease KB; Greenberg SJ; Longo DL

Metabolism Branch, National Cancer Institute, Bethesda, MD 20892.

Blood; 72(5):1805-16 1988 ISSN 0006-4971 Journal Code: A8G

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/38

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00687973 88320464 MEDL/88320464

Blockade of the interleukin-2 receptor by anti-Tac antibody inhibits the generation of antigen-nonspecific suppressor T cells in vitro. Oh-Ishi T; Goldman CK; Misiti J; Waldmann TA

Metabolism Branch, National Cancer Institute, Bethesda, MD 20892. Proc Natl Acad Sci U S A; 85(17):6478-82 1988 ISSN 0027-8424

Journal Code: PV3
Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/39

DIALOG(R)File 159:Cancerlit

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00682917 88285731 MEDL/88285731

Lateral diffusion measurements give evidence for association of the ${\bf Tac}$ peptide of the IL-2 receptor with the T27 peptide in the plasma membrane of HUT-102-B2 T cells.

Edidin M; Aszalos A; Damjanovich S; Waldmann TA

Department of Biology, Johns Hopkins University, Baltimore, MD 21218.

J Immunol; 141(4):1206-10 1988 ISSN 0022-1767 Journal Code: IEB

J Immunol; 141(4):1206-10 1988 ISSN 0022-1767 Journal Code: IFB Contract/Grant No.: AI-14584, SS, AI, SS, NIAID; AI-19814, SS, AI, SS, NIAID

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/40

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00665767 88151572 MEDL/88151572

Prolongation of primate renal allografts with anti-Tac monoclonal antibody.

Reed MH; Shapiro ME; Strom TB; Milford EL; Carpenter CB; Letvin NL; Waldmann TA; Kirkman RL

Curr Surg; 45(1):28-30 1988 ISSN 0149-7944 Journal Code: DWP

Contract/Grant No.: NO1-AI-52587, SS, AI, SS, NIAID

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/41

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00647210 88082577 MEDL/88082577

The role of the multichain IL-2 receptor complex in the control of normal and malignant T-cell proliferation.

Waldmann TA

National Cancer Institute, Bethesda, MD 20892.

Environ Health Perspect; 75:11-5 1987 ISSN 0091-6765 Journal Code: EIO Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL

5/3/42

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00641017 88041084 MEDL/88041084

Flow cytometric resonance energy transfer measurements support the association of a 95-kDa peptide termed T27 with the 55-kDa ${f Tac}$ peptide.

Szollosi J; Damjanovich S; Goldman CK; Fulwyler MJ; Aszalos AA; Goldstein G; Rao P; Talle MA; Waldmann ${\tt TA}$

Department of Biophysics, Medical University School of Debrecen, Hungary. Proc Natl Acad Sci U S A; 84(20):7246-50 1987 ISSN 0027-8424 Journal Code: PV3

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/43

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00634465 87322693 MEDL/87322693

The interleukin-2 receptor on normal and malignant lymphocytes.

Waldmann TA

Adv Exp Med Biol; 213:129-37 1987 ISSN 0065-2598 Journal Code: 2LU

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/44

DIALOG(R) File 159: Cancerlit

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00625742 87260992 MEDL/87260992

The p75 peptide is the receptor for interleukin 2 expressed on large granular lymphocytes and is responsible for the interleukin 2 activation of these cells.

Tsudo M; Goldman CK; Bongiovanni KF; Chan WC; Winton EF; Yagita M; Grimm EA; Waldmann TA

Proc Natl Acad Sci U S A; 84(15):5394-8 1987 ISSN 0027-8424

Journal Code: PV3 Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/45

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00621664 87231976 MEDL/87231976

Contribution of a p75 interleukin 2 binding peptide to a high-affinity interleukin 2 receptor complex.

Tsudo M; Kozak RW; Goldman CK; Waldmann TA

Proc Natl Acad Sci U S A; 84(12):4215-8 1987 ISSN 0027-8424

Journal Code: PV3
Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/46

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00617179 87196414 MEDL/87196414

Expression of functional IL 2 receptors by lipopolysaccharide and interferon-gamma stimulated human monocytes.

Holter W; Goldman CK; Casabo L; Nelson DL; Greene WC; Waldmann TA J Immunol; 138(9):2917-22 1987 ISSN 0022-1767 Journal Code: IFB Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/47

DIALOG(R) File 159: Cancerlit

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00596064 87092314 MEDL/87092314

Demonstration of a non-Tac peptide that binds interleukin 2: a potential participant in a multichain interleukin 2 receptor complex.

Tsudo M; Kozak RW; Goldman CK; Waldmann TA

Proc Natl Acad Sci U S A; 83(24):9694-8 1986 ISSN 0027-8424

Journal Code: PV3

Contract/Grant No.: 1F32 CA07974-01, SS, CA, SS, NCI

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/48

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00583605 87002626 MEDL/87002626

The interleukin-2 receptor on malignant cells: a target for diagnosis and therapy.

Waldmann TA

Cell Immunol; 99(1):53-60 1986 ISSN 0008-8749 Journal Code: CQ9

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/49

DIALOG(R)File 159:Cancerlit

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00581940 86315189 MEDL/86315189

The arrangement of immunoglobulin, T cell antigen receptor, and interleukin 2 receptor genes in human lymphoid neoplasms.

Waldmann TA; Korsmeyer SJ; Greene WC

Symp Fundam Cancer Res; 38:63-78 1986 ISSN 0190-1214 Journal Code: SFC

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/50

DIALOG(R) File 159: Cancerlit

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00567958 86217733 MEDL/86217733

Selective killing of human T-lymphotropic virus-I infected leukemic T-cells by monoclonal anti-interleukin 2 receptor antibody-ricin A chain conjugates: potentiation by ammonium chloride and monensin.

Kronke M; Schlick E; Waldmann TA; Vitetta ES; Greene WC

Cancer Res; 46(7):3295-8 1986 ISSN 0008-5472 Journal Code: CNF

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/51

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00559918 86149324 MEDL/86149324

Only high-affinity receptors for interleukin 2 mediate internalization of ligand.

Weissman AM; Harford JB; Svetlik PB; Leonard WL; Depper JM; Waldmann TA; Greene WC; Klausner RD

Proc Natl Acad Sci U S A; 83(5):1463-6 1986 ISSN 0027-8424

Journal Code: PV3 Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/52

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00555671 86094413 MEDL/86094413

Bismuth-212-labeled anti-Tac monoclonal antibody:

alpha-particle-emitting radionuclides as modalities for radioimmunotherapy.

Kozak RW; Atcher RW; Gansow OA; Friedman AM; Hines JJ; Waldmann TA Proc Natl Acad Sci U S A; 83(2):474-8 1986 ISSN 0027-8424

Journal Code: PV3

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/53

DIALOG(R) File 159: Cancerlit

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00554462 88640034 ICDB/88640034

ROLE AND MOLECULAR BIOLOGY OF THE INTERLEUKIN-2-INTERLEUKIN-2 RECEPTOR SYSTEM IN HEALTH AND DISEASE

Waldmann TA; Kozak RW; Tsudo M; Oh-ishi T; Bongiovanni KF; Goldman

Metabolism Branch, NCI, Bethesda, MD 20892

Non-serial; Progress in Immunology VI. Sixth International Congress of Immunology. Cinader B, Miller RG, eds. New York, Academic Press, p. 553-62, 1986.: 1986

Languages: ENGLISH

Document Type: MEETING PAPER

5/3/54

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00522623 85254471 MEDL/85254471

Isolation and expression of complementary DNAs encoding the human interleukin 2 receptor.

Greene WC; Depper JM; Crabtree GR; Rudikoff S; Pumphrey J; Robb RJ; Kronke M; Svetlik P; Peffer NJ; Waldmann TA; et al

Cancer Res; 45(9 Suppl):4563s-4567s 1985 ISSN 0008-5472 Journal Code: CNF

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/55

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00522622 85254470 MEDL/85254470

Interleukin 2 receptor (**Tac** antigen) expression in HTLV-I-associated adult T-cell leukemia.

Waldmann TA; Longo DL; Leonard WJ; Depper JM; Thompson CB; Kronke M; Goldman CK; Sharrow S; Bongiovanni K; Greene WC

Cancer Res; 45(9 Suppl):4559s-4562s 1985 ISSN 0008-5472 Journal Code:

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/56

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00514976 85200298 MEDL/85200298

Adult T cell leukemia: a potential target for ricin A chain immunotoxins. Kronke M; Depper JM; Leonard WJ; Vitetta ES; Waldmann TA; Greene WC Blood; 65(6):1416-21 1985 ISSN 0006-4971 Journal Code: A8G

Contract/Grant No.: CA-28149, SS, CA, SS, NCI

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/57

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00499850 86111567 MEDL/86111567

Interleukin-2 receptor expression in retrovirus associated adult T-cell leukemia.

Waldmann TA; Leonard WJ; Depper JM; Kronke M; Goldman CK; Oh T; Greene WC

Princess Takamatsu Symp; 15:259-68 1984 Journal Code: HHI

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/58

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00489169 85061621 MEDL/85061621

Human interleukin-2 promotes proliferation of activated B cells via surface receptors similar to those of activated T cells.

Mingari MC; Gerosa F; Carra G; Accolla RS; Moretta A; Zubler RH; Waldmann TA; Moretta L

Nature; 312(5995):641-3 1984 ISSN 0028-0836 Journal Code: NSC

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/59

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00484993 85032518 MEDL/85032518

Expression of interleukin 2 receptors on activated human B cells. Waldmann TA; Goldman CK; Robb RJ; Depper JM; Leonard WJ; Sharrow SO; Bongiovanni KF; Korsmeyer SJ; Greene WC

J Exp Med; 160(5):1450-66 1984 ISSN 0022-1007 Journal Code: I2V

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/60

DIALOG(R)File 159:Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00484844 85031785 MEDL/85031785

Regulation of interleukin 2 receptor expression: effects of phorbol diester, phospholipase C, and reexposure to lectin or antigen.

Depper JM; Leonard WJ; Kronke M; Noguchi PD; Cunningham RE; Waldmann TA; Greene WC

J Immunol; 133(6):3054-61 1984 ISSN 0022-1767 Journal Code: IFB

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/61

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00478032 84289961 MEDL/84289961

Pseudomonas exotoxin-anti-TAC . Cell-specific immunotoxin active

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against cells expressing the human T cell growth factor receptor.
  FitzGerald DJ; Waldmann TA; Willingham MC; Pastan I
  J Clin Invest; 74(3):966-71 1984 ISSN 0021-9738 Journal Code: HS7
 Languages: ENGLISH
  Document Type: JOURNAL ARTICLE
 5/3/62
DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
00471239
           84241073
                     MEDL/84241073
   Phorbol diester induces expression of Tac antigen on human acute T
lymphocytic leukemic cells.
  Greene WC; Robb RJ; Depper JM; Leonard WJ; Drogula C; Svetlik PB;
Wong-Staal F; Gallo RC; Waldmann TA
  J Immunol; 133(2):1042-7 1984 ISSN 0022-1767 Journal Code: IFB
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE
 5/3/63
DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
           84213140
                     MEDL/84213140
  Functional and phenotypic comparison of human T cell leukemia/lymphoma
virus positive adult T cell leukemia with human T cell leukemia/lymphoma
virus negative Sezary leukemia, and their distinction using anti-Tac.
Monoclonal antibody identifying the human receptor for T cell growth
factor.
  Waldmann TA; Greene WC; Sarin PS; Saxinger C; Blayney DW; Blattner
WA; Goldman CK; Bongiovanni K; Sharrow S; Depper JM; et al
  J Clin Invest; 73(6):1711-8 1984 ISSN 0021-9738 Journal Code: HS7
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE
 5/3/64
DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
00441769
           84070747
                     MEDL/84070747
  Characterization of the human receptor for T-cell growth factor.
  Leonard WJ; Depper JM; Robb RJ; Waldmann TA; Greene WC
  Proc Natl Acad Sci U S A; 80(22):6957-61 1983 ISSN 0027-8424
Journal Code: PV3
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE
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5/3/65

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00437919 84039776 MEDL/84039776

Association of human T-cell leukaemia/lymphoma virus with the ${\bf Tac}$ antigen marker for the human T-cell growth factor receptor.

Lando Z; Sarin P; Megson M; Greene WC; Waldman TA; Gallo RC; Broder

Nature; 305(5936):733-6 1983 ISSN 0028-0836 Journal Code: NSC

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

• 00428272 83273653 MEDL/83273653

Rearrangement and expression of immunoglobulin genes and expression of **Tac** antigen in hairy cell leukemia.

Korsmeyer SJ; Greene WC; Cossman J; Hsu SM; Jensen JP; Neckers LM; Marshall SL; Bakhshi A; Depper JM; Leonard WJ; Jaffe ES; Waldmann TA

Proc Natl Acad Sci U S A; 80(14):4522-6 1983 ISSN 0027-8424

Journal Code: PV3
Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/67

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00357793 83601683 ICDB/83601683

A MONOCLONAL ANTIBODY THAT APPEARS TO RECOGNIZE THE RECEPTOR FOR HUMAN T-CELL GROWTH FACTOR; PARTIAL CHARACTERIZATION OF THE RECEPTOR

Leonard WJ; Depper JM; Uchiyama T; Smith KA; Waldmann TA; Greene WC

Metabolism Branch, NCI, NIH, Bethesda, MD, 20205

Nature; 300(5889):267-269 1982

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/68

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00304399 81197499 MEDL/81197499

Activation of leukemic pro-suppressor cells to become suppressor-effector cells. Influence of cooperating normal T cells.

Broder S; Uchiyama T; Muul LM; Goldman C; Sharrow S; Poplack DG;

Waldmann TA

N Engl J Med; 304(23):1382-7 1981 ISSN 0028-4793 Journal Code: NOW

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

5/3/69

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00299993 81143481 MEDL/81143481

A monoclonal antibody (anti-Tac) reactive with activated and functionally mature human T cells. I. Production of anti-Tac monoclonal antibody and distribution of Tac (+) cells.

Uchiyama T; Broder S; Waldmann TA

J Immunol; 126(4):1393-7 1981 ISSN 0022-1767 Journal Code: IFB

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

11/3/2

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

92034763 MEDL/92034763 00889575

Biological monitoring of low-dose interleukin 2 in humans: soluble

interleukin 2 receptors, cytokines, and cell surface phenotypes.

Hanninen EL; Korfer A; Hadam M; Schneekloth C; Dallmann I; Menzel T;
Kirchner H; Poliwoda H; Atzpodien J

Department of Hematology and Oncology, Medizinische Hochschule Hanover,

Cancer Res; 51(23 Pt 1):6312-6 1991 ISSN 0008-5472 Journal Code: CNF Languages: ENGLISH

s tac and lymphoma? 1376 TAC 79436 LYMPHOMA? 168 TAC AND LYMPHOMA? S6 ? s s6 and py=1993 168 S6 80574 PY=1993 12 S6 AND PY=1993 **S7** ? rd s7 ...completed examining records 12 RD S7 (unique items) ? s s6 and py=1994168 S6 83740 PY=1994 10 S6 AND PY=1994 S9 ? t s7/3/all 7/3/1 DIALOG(R) File 159: Cancerlit (c) format only 2000 Dialog Corporation. All rts. reserv. 94036762 MEDL/94036762 01040145 Alternative splicing of the p53 tumor suppressor gene in the Molt-4 T-lymphoblastic leukemia cell line. Chow VT; Quek HH; Tock EP Department of Microbiology, Faculty of Medicine, National University of Singapore, Kent Ridge. Cancer Lett; 73(2-3):141-8 1993 ISSN 0304-3835 Journal Code: CMX Languages: ENGLISH Document Type: JOURNAL ARTICLE 7/3/2 DIALOG(R) File 159: Cancerlit (c) format only 2000 Dialog Corporation. All rts. reserv. 01034122 94003208 MEDL/94003208 The interleukin-2 receptor: a target for monoclonal antibody treatment of human T-cell lymphotrophic virus I-induced adult T-cell leukemia. Waldmann TA; White JD; Goldman CK; Top L; Grant A; Bamford R; Roessler E; Horak ID; Zaknoen S; Kasten-Sportes C; et al Metabolism Branch and Laboratory of Pathology, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892. Blood; 82(6):1701-12 1993 ISSN 0006-4971 Journal Code: A8G Languages: ENGLISH Document Type: JOURNAL ARTICLE 7/3/3 DIALOG(R) File 159: Cancerlit (c) format only 2000 Dialog Corporation. All rts. reserv.

MEDL/93368196

1992 Stohlman Memorial Lecture: targeting the IL-2 receptor.

93368196

Waldmann TA

Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892.

Leukemia; 7 Suppl 2:S151-6 1993 ISSN 0887-6924 Journal Code: LEU

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

7/3/4

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01021107 93318559 MEDL/93318559

Ki-1-positive large cell anaplastic **lymphoma** diagnosed by urinary cytology. A case report.

Tanaka T; Yoshimi N; Sawada K; Takami T; Sugie S; Etori F; Kachi H; Mori

Department of Pathology, Gifu University School of Medicine, Japan. Acta Cytol; 37(4):520-4 1993 ISSN 0001-5547 Journal Code: 0LI

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

7/3/5

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01019871 93311229 MEDL/93311229

Clinicopathologic, enzyme and histochemical studies of centrocytic (mantle cell) lymphoma: comparison with other types of low-grade B cell lymphoma based on the updated Kiel classification.

Takeshita M; Masuda Y; Sumiyoshi Y; Ohshima K; Kikuchi M; Kimura N; Okamura T; Nishimura J; Kozuru M

Department of Pathology, School of Medicine, Fukuoka University, Japan. Acta Pathol Jpn; 43(5):244-52 1993 ISSN 0001-6632 Journal Code: 1NE Languages: ENGLISH

Document Type: JOURNAL ARTICLE

7/3/6

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01004177 93218273 MEDL/93218273

Cytotoxic activities of recombinant immunotoxins composed of Pseudomonas toxin or diphtheria toxin toward lymphocytes from patients with adult T-cell leukemia.

Kreitman RJ; Chaudhary VK; Waldmann TA; Hanchard B; Cranston B; FitzGerald DJ; Pastan I

Laboratory of Molecular Biology, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892.

Leukemia; 7(4):553-62 1993 ISSN 0887-6924 Journal Code: LEU

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

7/3/7

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00991310 95606931 ICDB/95606931

Lymphokine receptors: a target for immunotherapy of lymphomas (Meeting abstract).

Waldmann TA

NCI, Bethesda, MD 20892

Non-serial; Fifth International Conference on Malignant Lymphoma, June 9-12, 1993, Lugano, Switzerland, p. 46, 1993.: 1993

Languages: ENGLISH

Document Type: MEETING ABSTRACTS

7/3/8

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00990752 95605905 ICDB/95605905

Progress in radioimmunotherapy (Meeting abstract).

Carrasquillo JA

Nuclear Medicine Department, Warren G. Magnuson Clinical Center, National Institutes of Health, Bethesda, MD 20892

Non-serial; International Congress of Radiation Oncology 1993. June 21-25, 1993, Kyoto, Japan, p. 149, 1993.: 1993

Languages: ENGLISH

Document Type: MEETING ABSTRACTS

7/3/9

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00986602 94690741 ICDB/94690741

Value of lymphography in the study of lymphomas]

Valor de la linfografia en el estudio de los linfomas.

Jimenez Cazorla AD

Universidad de Sevilla, Spain

Diss Abstr Int [C]; 54(1):245 1993 ISSN 0419-4217

Languages: SPANISH
Document Type: THESIS

7/3/10

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00985732 93694384 ICDB/93694384

Recombinant toxins for cancer treatment (Meeting abstract).

Pastan I; FitzGerald DJ

Lab. of Molecular Biology, NCI, Bethesda, MD 20892

Proc Annu Meet Am Assoc Cancer Res; 34:622-3 1993 ISSN 0197-016X

Languages: ENGLISH

Document Type: MEETING ABSTRACTS

7/3/11

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00985141 93693792 ICDB/93693792

Towards optimal anti-Tac antibody dosing in the presence of antigenemia in adult T-cell leukemia-lymphoma (Meeting abstract).

Junghans RP; Goldman CK; Carrasquillo J; Reynolds J; Nelson DL; Waldmann

New England Deaconess Hosp., Boston, MA 02215

Proc Annu Meet Am Assoc Cancer Res; 34:A2849 1993 ISSN 0197-016X

Languages: ENGLISH

Document Type: MEETING ABSTRACTS

7/3/12

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00983622 93692207 ICDB/93692207

Pilot Phase II study of a DAB486IL-2 fusion toxin in cutaneous T-cell lymphoma (Meeting abstract).

Foss F; Borkowski T; Tompkins A; Gilliom M; Cooper M; Udey M; Stetler-Stevenson M; Jaffe E; Woodworth T; Sausville E

NCI, Bethesda, MD

Proc Annu Meet Am Assoc Cancer Res; 34:A1236 1993 ISSN 0197-016X

Languages: ENGLISH

Document Type: MEETING ABSTRACTS

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9/3/1

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01138480 95182475 MEDL/95182475

Experimental study on anti-tumor effect of splenocytes induced by anti-CD3 McAb, PHA and IL-2.

Shen GX; Wang XL; Zhu HF; Zhang Y; Shao JF

Department of Immunology, Tongji Medical University, Wuhan.

J Tongji Med Univ; 14(1):12-5 1994 ISSN 0257-716X Journal Code: KAJ

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

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DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01109205 94362224 MEDL/94362224

Chimeric fusion protein toxin DAB486IL-2 in advanced mycosis fungoides and the Sezary syndrome: correlation of activity and interleukin-2 receptor expression in a phase II study.

Foss FM; Borkowski TA; Gilliom M; Stetler-Stevenson M; Jaffe ES; Figg WD; Tompkins A; Bastian A; Nylen P; Woodworth T; et al

US Naval Hospital, Bethesda, MD.

Blood; 84(6):1765-74 1994 ISSN 0006-4971 Journal Code: A8G

Languages: ENGLISH

Document Type: CLINICAL TRIAL; CLINICAL TRIAL, PHASE II; JOURNAL ARTICLE

9/3/3

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01102721 94325831 MEDL/94325831

Down-regulation of CD3 antigen on adult T cell leukemia cells.

Maeda Y; Matsuda M; Irimajiri K; Horiuchi A

Third Department of Internal Medicine, Kinki University School of Medicine, Osaka, Japan.

Leuk Lymphoma; 13(3-4):249-56 1994 ISSN 1042-8194 Journal Code: BNQ

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

9/3/4

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01102643 94325579 MEDL/94325579

Treatment of acute graft-versus-host disease with humanized anti-Tac: an antibody that binds to the interleukin-2 receptor.

Anasetti C; Hansen JA; Waldmann TA; Appelbaum FR; Davis J; Deeg HJ; Doney K; Martin PJ; Nash R; Storb R; et al

Clinical Research Division, Fred Hutchinson Cancer Research Center 98104. Blood; 84(4):1320-7 1994 ISSN 0006-4971 Journal Code: A8G

Contract/Grant No.: CA 18029, SS, CA, SS, NCI; CA 18221, SS, CA, SS, NCI;

HL36444, SS, HL, SS, NHLBI; +

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

9/3/5

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01101755 94320068 MEDL/94320068

Radioimmunotherapy of nude mice bearing a human interleukin 2 receptor alpha-expressing lymphoma utilizing the alpha-emitting radionuclide-conjugated monoclonal antibody 212Bi-anti-Tac.

Hartmann F; Horak EM; Garmestani K; Wu C; Brechbiel MW; Kozak RW; Tso J; Kosteiny SA; Gansow OA; Nelson DL; et al

Metabolism Branch, National Cancer Institute, NIH, Bethesda, Maryland

Cancer Res; 54(16):4362-70 1994 ISSN 0008-5472 Journal Code: CNF

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

9/3/6

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01097860 94297519 MEDL/94297519

Recombinant single-chain immunotoxins against T and B cell leukemias.

Kreitman RJ; Pastan I

Laboratory of Molecular Biology, National Cancer Institute, National Institutes of Health, Bethesda, Maryland 20892.

Leuk Lymphoma; 13(1-2):1-10 1994 ISSN 1042-8194 Journal Code: BNQ

Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, ACADEMIC

9/3/7

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

01092742 94266289 MEDL/94266289

Anti-IL-2 receptor monoclonal antibody (anti-Tac) treatment of T-cell lymphoma.

Waldmann TA

Metabolism Branch, National Cancer Institute, Bethesda, Maryland. Important Adv Oncol; :131-41 1994 ISSN 0883-5896 Journal Code: GG9 Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, ACADEMIC

9/3/8

DIALOG(R) File 159: Cancerlit

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01089877 94249675 MEDL/94249675

Differential expression of interleukin-2 receptors (alpha and beta chain) in mature lymphoid neoplasms.

Nakase K; Kita K; Nasu K; Ueda T; Tanaka I; Shirakawa S; Tsudo M Department of Internal Medicine, Yamada Red Cross Hospital, Misono, Japan.

Am J Hematol; 46(3):179-83 1994 ISSN 0361-8609 Journal Code: 3H4

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

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DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
           94226936
                    MEDL/94226936
01086036
  Lymphokine receptors: a target for immunotherapy of lymphomas.
  Waldmann TA
  Metabolism Branch, National Cancer Institute, National Institutes of
Health, Bethesda, Maryland.
  Ann Oncol; 5 Suppl 1:13-7 1994 ISSN 0923-7534
                                                   Journal Code: AYF
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL
 9/3/10
DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
01071386
           96605013
                      ICDB/96605013
  Effectiveness of bcl-2 antisense oligodeoxy-nucleotides (AS-ODN) against
        follicular small-cleaved cell lymphoma (FSCCL)-SCID mice
xenograft model (Meeting abstract).
  Abubakr YA; Mohammad R; Maki A; Dan M; Du W; Smith MR; Al-Katib A
  Division of Hematology/Oncology and Department of Pathology, Wayne State
University, Detroit, MI
  Blood; 84(10, Suppl 1):374a 1994 ISSN 0903-1936
  Languages: ENGLISH
  Document Type: MEETING ABSTRACTS
? ds
        Items
                Description
Set
                E1,E19
S1
          254
                S1 AND TAC
S2
           72
               S2 AND PY=1993
S3
            6
                S2 AND PY=1994
S4
           4
                RD S2 (unique items)
S5
           69
                TAC AND LYMPHOMA?
S6
          168
                S6 AND PY=1993
S7
           12
                RD S7 (unique items)
S8
           12
                S6 AND PY=1994
           10
S9
? s s6 and soluble (10n) (IL(w) 2 or IL(w) 2R)
             168 S6
           20463 SOLUBLE
           61351
                  IL
          499237
                  2
           21778 IL(W)2
           61351 IL
            3103
                  2R
            2251
                  IL(W)2R
             826 SOLUBLE(10N)(IL(W)2 OR IL(W)2R)
               5 S6 AND SOLUBLE (10N) (IL(W) 2 OR IL(W) 2R)
     S10
? rd s10
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...completed examining records

5 RD S10 (unique items)

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s soluble (30n) (IL(w)2 or IL(w)2R or IL(w)2(w) receptor?) and tac
           20463 SOLUBLE
           61351
                 IL
          499237
           21778 IL(W)2
           61351
                 IL
            3103
                 2R
            2251 IL(W)2R
           61351
                 IL
          499237
          122702 RECEPTOR?
            3467 IL(W) 2(W) RECEPTOR?
            1164 SOLUBLE (30N) ((IL(W)2 OR IL(W)2R) OR IL(W)2(W)RECEPTOR?)
           1376 TAC
     S12
             43
                 SOLUBLE (30N) (IL (W) 2 OR IL (W) 2R OR IL (W) 2 (W) RECEPTOR?) AND
                  TAC
? rd s12
...completed examining records
     S13
             43 RD S12 (unique items)
? t s13/3/all
13/3/1
DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
01095932
          94286643
                    MEDL/94286643
  Targeting human IL-2 receptors for diagnosis and therapy.
  Nelson DL; Kurman CC
  Immunophysiology Section, National Cancer Institute, National Institutes
of Health, Bethesda, Maryland 20892.
  Proc Soc Exp Biol Med; 206(3):309-11 1994 ISSN 0037-9727
Journal Code: PXZ
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL
13/3/2
DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
01075134
          94139113
                    MEDL/94139113
   Follow up of soluble IL-2 receptor level in
metastatic malignant melanoma patients treated by chemoimmunotherapy.
  Soubrane C; Mouawad R; Ichen M; Suissa J; Borel C; Vuillemin E;
Benhammouda A; Bizzari JP; Weil M; Khayat D
  Department of Medical Oncology, Salpetriere Hospital, Paris, France.
  Clin Exp Immunol; 95(2):232-6 1994 ISSN 0009-9104 Journal Code: DD7
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE
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DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
           94118567 MEDL/94118567
01073616
  Does IL-2 receptor expression and secretion in chronic B-cell leukemia
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have a role in down-regulation of the immune system?

Burton J; Kay NE

Metabolism Branch, National Institute of Health, Bethesda, MD. Leukemia; 8(1):92-6 1994 ISSN 0887-6924 Journal Code: LEU

Languages: ENGLISH

Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL

13/3/4

DIALOG(R) File 159: Cancerlit

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01049259 94092956 MEDL/94092956

Human T lymphocyte activation by pokeweed mitogen induces production of TNF-alpha and GM-CSF and helper signaling by IL-1 and IL-6 results in IL-2-dependent T cell growth.

Wallays G; Ceuppens JL

Department of Medicine and Pathophysiology, University of Leuven, Belgium.

Eur Cytokine Netw; 4(4):269-77 1993 ISSN 1148-5493 Journal Code: A56

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

13/3/5

DIALOG(R) File 159: Cancerlit

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01034189 94003317 MEDL/94003317

A model of in vivo cell proliferation of adult T-cell leukemia.

Kondo A; Imada K; Hattori T; Yamabe H; Tanaka T; Miyasaka M; Okuma M;
Uchiyama T

First Department of Internal Medicine and Laboratory of Anatomic Pathology, Kyoto University, Japan.

Blood; 82(8):2501-9 1993 ISSN 0006-4971 Journal Code: A8G

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

13/3/6

DIALOG(R) File 159: Cancerlit

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00966791 93077183 MEDL/93077183

Chemo-immunotherapy in patients with metastatic melanoma using sequential treatment with dacarbazine and recombinant human interleukin-2: evaluation of hematologic and immunologic parameters and correlation with clinical response.

Isacson R; Kedar E; Barak V; Gazit Z; Yurim O; Kalichman I; Ben-Bassat H; Biran S; Schlesinger M; Franks CR; et al

Department of Oncology, Hadassah University Hospital, Jerusalem, Israel. Immunol Lett; 33(2):127-34 1992 ISSN 0165-2478 Journal Code: GIH

Languages: ENGLISH

Document Type: CLINICAL TRIAL; CLINICAL TRIAL, PHASE II; JOURNAL ARTICLE; MULTICENTER STUDY

13/3/7

DIALOG(R) File 159: Cancerlit

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00950624 92388287 MEDL/92388287

Membrane-based receptor affinity chromatography.

Nachman M; Azad AR; Bailon P

Protein Biochemistry Department, Roche Research Center, Hoffmann-La Roche Inc., Nutley, NJ 07110.

J Chromatogr; 597(1-2):155-66 1992 ISSN 0021-9673 Journal Code: HQF

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

1)

13/3/8

DIALOG(R) File 159: Cancerlit

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00930410 92256201 MEDL/92256201

Diagnostic and clinical importance of interleukin-2 receptor alpha chain expression on non-T-cell acute leukaemia cells.

Nakase K; Kita K; Otsuji A; Anazawa H; Hoshino K; Sekine T; Shirakawa S; Tanaka I; Nasu K; Tsutani H; et al

Second Department of Internal Medicine, Faculty of Medicine, Mie University, Tsu, Japan.

Br J Haematol; 80(3):317-26 1992 ISSN 0007-1048 Journal Code: AXC

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

13/3/9

DIALOG(R) File 159: Cancerlit

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00924210 92208969 MEDL/92208969

Highly concentrated urine-purified **Tac** peptide fails to inhibit IL-2-dependent cell proliferation in vitro.

Pizzolo G; Vincenzi C; Vinante F; Rigo A; Veneri D; Chilosi M; Dusi S; Poli G; Zambello R; Semenzato G; et al

Cattedra di Ematologia, Istituto di Patologia Generale, Verona, Italy. Cell Immunol; 141(1):253-9 1992 ISSN 0008-8749 Journal Code: CQ9

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

13/3/10

DIALOG(R) File 159: Cancerlit

(c) format only 2000 Dialog Corporation. All rts. reserv.

00923445 92201878 MEDL/92201878

The monocyte interleukin-2 receptor light chain: production of cell-associated and soluble interleukin-2 receptor by monocytes.

Kniep EM; Strelow I; Lohmann-Matthes ML

Abteilung fur Immunobiologie, Fraunhofer Institut fur Toxikologie, Hannover, Germany.

Immunology; 75(2):299-304 1992 ISSN 0019-2805 Journal Code: GH7

Languages: ENGLISH

Document Type: JOURNAL ARTICLE

13/3/11

DIALOG(R) File 159: Cancerlit

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00914263 93689708 ICDB/93689708

Combination of rIFN alpha-2a and IL-2 in melanoma patients: what are effective doses? (Meeting abstract).

Castello G; Comella P; Napolitano M; Manzo T; Leonardi E; Galati MG; Casaretti R; Daponte A; Comella G

Natl. Cancer Inst. of Naples, Naples, Italy Ann Oncol; 3(Suppl 5):141 1992 ISSN 0923-7534

Languages: ENGLISH

Document Type: MEETING ABSTRACTS; CLINICAL TRIAL

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13/3/12
DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
                    MEDL/92136003
          92136003
 The cytokine receptor superfamily.
 Kaczmarski RS; Mufti GJ
 Department of Haematological Medicine, King's College School of Medicine
and Dentistry, London, UK.
 Blood Rev; 5(3):193-203 1991 ISSN 0268-960X
                                                Journal Code: BLR
 Languages: ENGLISH
 Document Type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL
13/3/13
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                    MEDL/92034763
00889575
          92034763
 Biological monitoring of low-dose interleukin 2 in humans: soluble
interleukin 2 receptors, cytokines, and cell surface phenotypes.
 Hanninen EL; Korfer A; Hadam M; Schneekloth C; Dallmann I; Menzel T;
Kirchner H; Poliwoda H; Atzpodien J
  Department of Hematology and Oncology, Medizinische Hochschule Hanover,
Germany.
 Cancer Res; 51(23 Pt 1):6312-6 1991 ISSN 0008-5472 Journal Code: CNF
 Languages: ENGLISH
 Document Type: JOURNAL ARTICLE
13/3/14
DIALOG(R) File 159: Cancerlit
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00881013
          91360958
                    MEDL/91360958
  [The interleukin-2 receptor (editorial)]
  Il recettore dell'interleukina-2.
 Marino P; Preatoni A
  Recenti Prog Med; 82(5):291-3 1991 ISSN 0034-1193 Journal Code: R1T
  Languages: ITALIAN
  Document Type: EDITORIAL English Abstract
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DIALOG(R) File 159: Cancerlit
(c) format only 2000 Dialog Corporation. All rts. reserv.
           91002416
                    MEDL/91002416
00819435
   Increased serum levels of soluble IL-2 receptor,
           CD8 molecules, and gamma-interferon in angioimmunoblastic
lymphadenopathy: possible pathogenetic role of immunoactivation mechanisms.
  Pizzolo G; Stein H; Josimovic-Alasevic O; Vinante F; Zanotti R; Chilosi M
; Feller AC; Diamantstein T
  Department of Hematology, Verona University, Italy.
  Br J Haematol; 75(4):485-8 1990 ISSN 0007-1048 Journal Code: AXC
  Languages: ENGLISH
  Document Type: JOURNAL ARTICLE
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The use of antibodies against the IL-2 receptor in transplantation

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0952791598003542

Waldmann_T_A O'Shea_J

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ISSN

0952-7915

Publisher

Current Biology

Year of publication Volume

1998

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2/3/53 (Item 4 from file: 399) DIALOG(R) File 399:CA SEARCH(R) (c) 2000 American Chemical Society. All rts. reserv.

111005761 CA: 111(1)5761u PATENT

Monoclonal antibody to Tac receptor for treatment of malignancy and autoimmune disorders in humans

INVENTOR (AUTHOR): Waldmann, T. A.

LOCATION: USA

ASSIGNEE: United States Dept. of Health and Human Services

PATENT: U.S. Pat. Appl. ; US 85707 A0 DATE: 880415

APPLICATION: US 85707 (870817)

PAGES: 18 pp. Avail. NTIS Order No. PAT-APPL-7-85707. CODEN: XAXXAV

2/3/52 (Item 3 from file: 399)
DIALOG(R)File 399:CA SEARCH(R)
(c) 2000 American Chemical Society. All rts. reserv.

117190132 CA: 117(19)190132z PATENT
Antibody composition directed against human or animal interleukin-2
(IL-2) receptor, and therapeutic uses thereof

INVENTOR(AUTHOR): Jacques, Yannick; Soulillou, Jean Paul; Audrain, Marie; Francois-Bendahou, Christine

LOCATION: Fr.

ASSIGNEE: Institut National de la Sante et de la Recherche Medicale (INSERM)

PATENT: PCT International; WO 9213886 Al DATE: 920820 APPLICATION: WO 92FR75 (920128) *FR 911102 (910131)

PAGES: 30 pp. CODEN: PIXXD2 LANGUAGE: French CLASS: C07K-015/00A DESIGNATED COUNTRIES: AU; CA; JP; KR; US DESIGNATED REGIONAL: AT; BE; CH

; DE; DK; ES; FR; GB; GR; IT; LU; MC; NL; SE

2/3/53 (Item 4 from file: 399)

2/3/29 n 1 from file: 73) DIALOG(R)Fi 3:EMBASE (c) 2000 El c Science B.V. All rts. reserv. 10828958 \SE No: 2000309668 /ivo blockade of CD25 (high-affinity interleukin 2 Successfu ells by administration of humanized anti-Tac receptor) o antibody to ents with psoriasis Krueger J Valters I.B.; Miyazawa M.; Gilleaudeau P.; Hakimi J.; Light S.; Sherr A :tlieb A.B. Dr. J.G. er, Laboratory Head, Rockefeller University, Lab. for Investiga Dermatology, 1230 York Ave, New York, NY 10021-6399 United States AUTHOR EM Journal o American Academy of Dermatology (J. AM. ACAD. DERMATOL.) (United Sta 2000, 43/3 (448-458) ISSN: 0190-9622 CODEN: JA Journal; Article DOCUMENT

SUMMARY LANGUAGE: ENGLISH

LANGUAGE:

[SH

10828958 EMBASE Successful in vivo receptor) on T cells antibody to patients Krueger J.G.; Walt S.; Sherr A.; Gottli Dr. J.G. Krueger, Investigative Derm States AUTHOR EMAIL: krue Journal of the Ame (United States) 200 ISS CODEN: JAADD DOCUMENT TYPE: Jou LANGUAGE: ENGLISH NUMBER OF REFERENC

Background: Dacliz (CD25) of the interl normal IL-2 binding T-cell growth, block be useful in treatir. Objective: Our purpo antibody were achiev lesions to saturate effect and safety of immunosuppressive dr psoriasis in two cer then 1 mg/kg at week blocked in vivo, flc CD3sup + T cells der of CD25sup + cells c Patients were follow and Severity Index s of CD25 in periphera while the dosing was began after 4 weeks, improvement. Patient score of less than 3 (P = .02). During th of the IL-2 receptor were calculated and study No significant study. Conclusion: W well-tolerated agent skin. Furthermore, i

10309668

1de of CD25 (high-affinity interleukin 2

1 ninistration of humanized anti-Tac

1 soriasis

3.; Miyazawa M.; Gilleaudeau P.; Hakimi J.; Light

1 tory Head, Rockefeller University, Lab. for

1 1230 York Ave, New York, NY 10021-6399 United

1 kvax.rockefeller.edu

1 cademy of Dermatology (J. AM. ACAD. DERMATOL.)

1 (448-458)

1 9622

1 rticle

1 RY LANGUAGE: ENGLISH

a humanized antibody to the alpha-subunit (IL-2) receptor that blocks receptor. Because IL-2 is a major stimulus for the IL-2 receptor could 1-mediated (autoimmune) diseases. to determine whether adequate concentrations of irculating blood and in psoriatic skin ceptors. We also intended to measure clinical .gent when used alone (without other psoriasis. Methods: Nineteen patients with ceived daclizumab at an initial dose of 2 mg/kg, 8, and 12. To determine whether CD25 was etric studies measured (1) expression of CD25 on om blood and (2) immunohistochemistry measures pretreatment and posttreatment biopsy specimens. ·linically with photographs and Psoriasis Area Results: This study showed a consistent blockade and tissue during the first 4 weeks of therapy 2 weeks. Variable desaturation of receptors correlated with a reversal in disease a pretreatment Psoriasis Area and Severity Index d a mean reduction in severity by 30% at 8 weeks eks of treatment, a 44.8% decrease in expression subunit was found. The absolute T-cell counts no significant changes during the course of the e events were produced by daclizumab during this fore conclude that daclizumab is a plocks CD25 expression in peripheral blood and we useful in treating psoriasis in some patients.

53 RD S1 (uni ns)
? s s2 and (soluble) (w) 2R or receptor?)

53 S2
326299 S0
256953 IL
21117 2R
8771 IL
1948284 RE
19857 SO
N) (IL(W) 2R OR RECEPTOR?)
S3 0 S2 DLUBLE) (5N) (IL(W) 2R OR RECEPTOR?)

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              25 S2 AND TAC
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DIALOG(R) File 5: Biosis Previews(R)
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13107907 BIOSIS NO.: 200100315056

Pretargeting radioimmunotherapy with the alpha emitter, BI-213, in the treatment of adult T cell leukemia (ATL).

AUTHOR: Zhang Meili(a); Yao Zhengsheng; Garmestani Kayhan; Axworthy Donald B; Mallett Robert W; Fritzberg Alan R; Brechbiel Martin W; Carrasquillo Jorge A; Waldmann Thomas A(a

AUTHOR ADDRESS: (a) Metabolism Branch, National Cancer Institute, NIH, Bethesda, MD**USA

JOURNAL: Blood 96 (11 Part 1):p721a November 16, 2000

MEDIUM: print

CONFERENCE/MEETING: 42nd Annual Meeting of the American Society of Hematology San Francisco, California, USA December 01-05, 2000

SPONSOR: American Society of Hematology

ISSN: 0006-4971

RECORD TYPE: Abstract LANGUAGE: English

SUMMARY LANGUAGE: English

ABSTRACT: Adult T-cell leukemia (ATL) consists of an overabundance of activated T cells which are characterized by the expression of CD25 or IL-2Ralpha on the cell surface. Presently, there is no accepted curative therapy for ATL. In this study, we investigated the use of a pretargeting technique with the alpha-emitter, Bi-213, to treat ATL. DOTA-biotin was labeled with Bi-213 at a specific activity of >1 mCi/mug. Humanized anti-Tac antibody (HAT), which recognizes CD25, was conjugated to streptavidin (SA) forming HAT-SA. A tumor model was established by i.p. injection of 20X106 tumor cells from human ATL-tumor bearing mice into NOD/SCID mice. The therapy study was performed at two weeks after the inoculation. Before therapy and every two weeks thereafter, serum soluble Tac (sIL-2Ralpha) levels were measured in order to monitor tumor growth. HAT-SA (140 mug) was used allowing pretargeting for 24 hours. Then, 100 mug of synthetic clearing agent (sCA), containing biotin and galactose arms, developed by NeoRx, was used to clear unbound conjugate from the circulation. Four hours later, 0.3 mug of Bi-213 labeled DOTA-biotin was given for therapy. Groups of 5 mice each were treated with 0, 50, 150, 250 or 350 muCi of Bi-213 and one group without any treatment served as a control. Complete blood count (CBC) was monitored for toxicity weekly. Results: The tumor growth was inhibited by using 250 or 350 muCi of Bi-213 labeled biotin in the pretargeting technique as indicated by the significant reduction of sIL-2Ralpha levels (p<0.005, t test) compared with the control group on day 28. Furthermore, there was a significant prolongation of the survival of the mice treated with either 250 or 350 muCi of Bi-213 as compared with the control group (p<0.02, rank test). The median survival duration of the control group was 52 days, whereas it increased to >75.6 and >67 days in 250 and 350 muCi Bi-213 treated groups, respectively. The CBC results revealed a dose dependant decrease of platelet and white blood cell count one week after therapy, with recovery three weeks later. Conclusion: Bi-213 labeled biotin used with the pretargeting technique proved to be an effective therapeutic for ATL without unacceptable toxicity.

3/7/2 (Item 2 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
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12862062 BIOSIS NO.: 200100069211

IL-2Ralpha-directed monoclonal antibodies provide effective therapy in a murine model of adult T-cell leukemia by a mechanism other than blockade of IL-2/IL-2Ralpha interaction.

AUTHOR: Phillips Kathleen E; Herring Bert; Wilson Latresia A; Rickford Marc S; Zhang Meili; Goldman Carolyn K; Tso J Yun; Waldmann Thomas A(a AUTHOR ADDRESS: (a) Metabolism Branch, National Cancer Institute, NIH, 10 Center Drive, Building 10, Room 4N-115, Bethesda, MD, 20892-1374**USA JOURNAL: Cancer Research 60 (24):p6977-6984 December 15, 2000

MEDIUM: print ISSN: 0008-5472

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

SUMMARY LANGUAGE: English

ABSTRACT: Adult T-cell leukemia (ATL) develops in a small proportion of human T-cell lymphotrophic virus-I infected individuals. The leukemia consists of an overabundance of activated T cells, which are characterized by the expression of CD25, or IL-2Ralpha, on their cell surface. Presently, there is not an accepted curative therapy for ATL. We developed an in vivo model of ATL in non-obese diabetic/severe combined immunodeficient (NOD/SCID) mice by introducing cells from an ATL patient (MET-1) into the mice. The leukemic cells proliferated in these mice that lack functional T, B, and natural killer (NK) cells. The MET-1 leukemic cells could be monitored by measurements of both serum soluble Tac (IL-2Ralpha) and soluble human beta2-microglobulin (beta2mu) by ELISA. The disease progressed to death in the mice after apprx4-6 weeks. The mice developed grossly enlarged spleens and a leukemia involving ATL cells that retained the phenotype and the T-cell receptor rearrangement and human T-cell lymphotrophic virus-I integration pattern of the patient's ATL leukemia cells. This model is of value for testing the efficacy of novel therapeutic agents for ATL. The administration of humanized anti-Tac (HAT), murine anti-Tac (MAT), and 7G7/B6, all of which target IL-2Ralpha, significantly delayed the progression of the leukemia and prolonged the survival of the tumor-bearing mice. In particular, HAT induced complete remissions in 4 of 19 mice and partial remissions in the remainder. It appears that the antibodies act by a mechanism that had not been anticipated. The prevailing view is that antibodies to the IL-2Ralpha receptor have their effective action by blocking the interaction of IL-2 with its growth factor receptor, thereby inducing cytokine deprivation apoptosis. However, although both HAT and MAT block the binding of IL-2 to IL-2Ralpha of the high affinity receptor, the 7G7/B6 monoclonal antibody binds to a different epitope on the IL-2Ralpha receptor, one that is not involved in IL-2 binding. This suggested that the antibodies provide an effective therapy by a mechanism other than induction of cytokine deprivation. In accord with this view, the MET-1 cells obtained from the spleens of leukemic mice did not produce IL-2, nor did they express IL-2 mRNA as assessed by reverse transcription-PCR. Another possible conventional mechanism of action involves complement-mediated killing. However, although MAT and 7G7/B6 fix rabbit complement, HAT does not do so. Furthermore, in the presence of NOD/SCID mouse serum, there was no complement-mediated lysis of MET-1 cells. In addition, the antibodies did not manifest antibody-dependent cellular cytotoxicity with NOD/SCID splenocytes that virtually lack NK cells as the effector cells as assessed in an in vitro chromium-release assay. However, in contrast to the efficacy of intact HAT, the F(ab')2 version of this antibody was not effective in prolonging the survival of mice injected with MET-1 ATL cells. In conclusion, in our murine model of ATL, monoclonal antibodies, HAT, MAT, and 7G7/B6, appear to delay progression of the leukemia by a mechanism of action that is different from the accepted mechanism of IL-2 deprivation leading to cell death. We consider two alternatives: the first, antibody-dependent cellular cytotoxicity mediated by FcRI- or FcRIII-expressing cells other than NK $\,$ cells, such as monocytes or polymorphonuclear leukocytes. The second alternative we consider involves direct induction of apoptosis by the anti-IL-2R antibodies in vivo. It has been shown that the IL-2R is a critical element in the peripheral self-tolerance T-cell suicide mechanism involved in the phenomenon of activation-induced cell death.

3/7/3 (Item 3 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
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12831477 BIOSIS NO.: 200100038626

Advances in interleukin 2 receptor targeted treatment.

AUTHOR: Morris John C(a); Waldmann Thomas A

AUTHOR ADDRESS: (a) Metabolism Branch, Division of Clinical Sciences, National Cancer Institute, NIH, 10 Center Drive, Bldg 10, Rm 4N115,

Bethesda, MD, 20892-1374: jmorris@mail.nih.gov**USA

JOURNAL: Annals of the Rheumatic Diseases 59 (Supplement 1):pi109-i114

November, 2000 MEDIUM: print ISSN: 0003-4967

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

SUMMARY LANGUAGE: English

ABSTRACT: T cell activation and cellular immune responses are modulated by interleukin 2 (IL2) through binding to its corresponding cell surface receptor. Three forms of the receptor are recognised based on IL2 binding affinity. The high affinity receptor is a heterotrimer composed of alpha, beta, and gammac-polypeptide chains. The 55 kDa alpha-chain also known as the Tac (T cell activation) antigen or CD-25 is a unique subunit of the high affinity IL2 receptor (IL2Ralpha). Resting T cells express few IL2Ralpha, however, when activated, the expression of ILR2alpha rapidly increases. The IL2Ralpha is shed from the cell surface and is measurable in the serum as a 45 kDa soluble form (s-Tac or s-IL2Ralpha). Serum concentrations of s-Tac can be used as a surrogate marker for T cell activation and IL2Ralpha expression. IL2Ralpha is over expressed by T cells in a number of autoimmune diseases, allograft rejection and a variety of lymphoid neoplasms. IL2 induced proliferation of T cells can be inhibited by the murine monoclonal antibody (anti-Tac) directed against the alpha-chain of the IL2R. Through molecular engineering, murine anti-Tac has been humanised reducing its immunogenicity without changing its specificity. Humanised anti-Tac (HAT) has been shown to reduce the incidence of renal and cardiac allograft rejection as well as decrease the severity of graft versus host disease in patients undergoing HLA matched allogeneic bone marrow transplantation. IL2Ralpha targeted treatment with radioimmunoconjugates of anti-Tac and immunotoxins has shown promise in the treatment of CD25 expressing lymphomas.

3/7/4 (Item 4 from file: 5)
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12482143 BIOSIS NO.: 200000235645

Phase I trial of recombinant immunotoxin anti-Tac(Fv)-PE38 (LMB-2) in patients with hematologic malignancies.

AUTHOR: Kreitman Robert J; Wilson Wyndham H; White Jeffrey D; Stetler-Stevenson Maryalice; Jaffe Elaine S; Giardina Steven; Waldmann Thomas A; Pastan Ira(a

AUTHOR ADDRESS: (a) Laboratory of Molecular Biology, National Cancer Institute, National Institute of Health, 9000 Rockville Pike, Building 37/4E16, Bethesda, MD, 20892**USA

JOURNAL: Journal of Clinical Oncology 18 (8):p1622-1636 April, 2000

ISSN: 0732-183X

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

SUMMARY LANGUAGE: English

ABSTRACT: Purpose: To evaluate the toxicity, pharmacokinetics, immunogenicity, and antitumor activity of anti-**Tac**(Fv)-PE38 (LMB-2), an anti-CD25 recombinant immunotoxin that contains an antibody Fv fragment fused to truncated Pseudomonas exotoxin. Patients and Methods: Patients with CD25+ hematologic malignancies for whom standard

and salvage therapies failed were treated with LMB-2 at dose levels that ranged from 2 to 63 mug/kg administered intravenously over 30 minutes on alternate days for three doses (QOD X 3). Results: LMB-2 was administered to 35 patients for a total of 59 cycles. Dose-limiting toxicity at the 63 mug/kg level was reversible and included transaminase elevations in one patient and diarrhea and cardiomyopathy in another. LMB-2 was well tolerated in nine patients at the maximum-tolerated dose (40 mug/kg QOD X 3); toxicity was transient and most commonly included transaminase elevations (eight patients) and fever (seven patients). Only six of 35 patients developed significant neutralizing antibodies after the first cycle. The median half-life was 4 hours. One hairy cell leukemia (HCL) patient achieved a complete remission, which is ongoing at 20 months. Seven partial responses were observed in cutaneous T-cell lymphoma (one patient), HCL (three patients), chronic lymphocytic leukemia (one patient), Hodgkin's disease (one patient), and adult T-cell leukemia (one patient). Responding patients had 2 to 5 log reductions of circulating malignant cells, improvement in skin lesions, and regression of lymphomatous masses and splenomegaly. All four patients with HCL responded to treatment. Conclusion: LMB-2 has clinical activity in CD25+ hematologic malignancies and is relatively nonimmunogenic. It is the first recombinant immunotoxin to induce major responses in cancer. LMB-2 and similar agents that target other cancer antigens merit further clinical development.

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12831477 BIOSIS NO.: 200100038626

Advances in interleukin 2 receptor targeted treatment.

AUTHOR: Morris John C(a); Waldmann Thomas A

AUTHOR ADDRESS: (a) Metabolism Branch, Division of Clinical Sciences, National Cancer Institute, NIH, 10 Center Drive, Bldg 10, Rm 4N115,

Bethesda, MD, 20892-1374: jmorris@mail.nih.gov**USA

JOURNAL: Annals of the Rheumatic Diseases 59 (Supplement 1):pil09-il14

November, 2000 MEDIUM: print ISSN: 0003-4967

DOCUMENT TYPE: Article RECORD TYPE: Abstract

LANGUAGE: English

SUMMARY LANGUAGE: English

ABSTRACT: T cell activation and cellular immune responses are modulated by interleukin 2 (IL2) through binding to its corresponding cell surface receptor. Three forms of the receptor are recognised based on IL2 binding affinity. The high affinity receptor is a heterotrimer composed of alpha, beta, and gammac-polypeptide chains. The 55 kDa alpha-chain also known as the Tac (T cell activation) antigen or CD-25 is a unique subunit of the high affinity IL2 receptor (IL2Ralpha). Resting T cells express few IL2Ralpha, however, when activated, the expression of ILR2alpha rapidly increases. The IL2Ralpha is shed from the cell surface and is measurable in the serum as a 45 kDa soluble form (s-Tac or s-IL2Ralpha). Serum concentrations of s-Tac can be used as a surrogate marker for T cell activation and IL2Ralpha expression. IL2Ralpha is over expressed by T cells in a number of autoimmune diseases, allograft rejection and a variety of lymphoid neoplasms. IL2 induced proliferation of T cells can be inhibited by the murine monoclonal antibody (anti-Tac) directed against the alpha-chain of the IL2R. Through molecular engineering, murine anti-Tac has been humanised reducing its immunogenicity without changing its specificity. Humanised anti-Tac (HAT) has been shown to reduce the incidence of renal and cardiac allograft rejection as well as decrease the severity of graft versus host disease in patients undergoing HLA matched allogeneic bone marrow transplantation. IL2Ralpha targeted treatment with radioimmunoconjugates of anti-Tac and immunotoxins has shown promise in the treatment of CD25 expressing lymphomas.

```
? s (tac)(10n)(antibod?)(30n)(radiolabel? or yttrium? or
90Y) (40n) (mg) (40n) (treat? or therap? or administ?)
Processing
Processing
Processing
            8865
                  TAC
         1537832 ANTIBOD?
           92472 RADIOLABEL?
           81886 YTTRIUM?
            1178 90Y
         1234664 MG
         4863111
                  TREAT?
         4576406
                  THERAP?
         2662304 ADMINIST?
                  (TAC) (10N) (ANTIBOD?) (30N) (RADIOLABEL? OR YTTRIUM? OR
                  90Y) (40N) (MG) (40N) (TREAT? OR THERAP? OR ADMINIST?)
? rd s1
...completed examining records
               3 RD S1 (unique items)
? t s2/7/all
           (Item 1 from file: 5)
DIALOG(R) File 5:Biosis Previews(R)
(c) 2000 BIOSIS. All rts. reserv.
           BIOSIS NO.: 200000211864
12458362
Pharmacokinetics of 111In- and 125I-labeled antiTac single-chain Fv
  recombinant immunotoxin.
AUTHOR: Kobayashi Hisataka; Kao Chih-Hao K; Kreitman Robert J; Le Nhat; Kim
  Meyoung-Kon; Brechbiel Martin W; Paik Chang H; Pastan Ira; Carrasquillo
  Jorge A(a)
AUTHOR ADDRESS: (a) Department of Nuclear Medicine, National Institutes of
  Health, 10 Center Dr., Bldg. 10, Rm. 1C496, Bethesda, MD, 20892-1180**USA
JOURNAL: Journal of Nuclear Medicine 41 (4):p755-762 April, 2000
ISSN: 0161-5505
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English
SUMMARY LANGUAGE: English
ABSTRACT: The use of immunotoxins for cancer therapy is an attractive
  strategy that exploits the targeting specificity of monoclonal
  antibodies and their fragments as well as the exquisite toxicity of
  the toxins. However, few studies of immunotoxins have evaluated their
  biodistribution in vivo. Previous studies have used 125I for tracing
  immunotoxin biodistribution in mice. Because the immunotoxin works only
  when it is internalized and because of known problems with quick
  dehalogenation after internalization of antibodies, we decided to
  use 111In, which has greater intracellular retention than iodine.
  Methods: To trace the in vivo pharmacokinetics of the immunotoxin in
  mice, we labeled the antiTac(Fv)-PE38 with 111In and compared it with
  125I-labeled antiTac(Fv)-PE38. We successfully labeled antiTac(Fv)-PE38
```

with 111In at up to $2.96 \, \mathrm{GBq/mg}$. A 3- to 4-fold decrease in cytotoxicity was observed for both radiolabeled preparations. We

into ATAC4 cells (Tac-positive) as well as their biodistribution

evaluated the internalization of 111In- and 125I-labeled antiTac(Fv)-PE38

and pharmacokinetics in vivo in mice. In addition, some mice receiving these reagents were co-infused with 30 mg L-lysine to inhibit renal accumulation. Results: Significantly more 111In- than 125I-labeled antiTac(Fv)-PE38 accumulated in the ATAC4 cells (20% versus 5% of initial surface-bound radioactivity; P < 0.001). In vivo, significantly more 111In- than 125I-labeled antiTac(Fv)-PE38 accumulated in the kidney (119 versus 31 percentage injected dose per gram (%ID/g); P < 0.001). The tumor accumulation of 111In-labeled antiTac(Fv)-PE38 at 96 h was 13-fold greater than that of 125I-labeled antiTac(Fv)-PE38 (1.4 versus 0.1 %ID/g; P < 0.001). No antiTac(Fv)-PE38 was excreted into the urine in its intact form unless lysine was co-infused. Co-injected lysine reduced the renal accumulation of 111In-labeled antiTac(Fv)-PE38 by 62%. Conclusion: We evaluated the biodistribution, pharmacokinetics, and catabolism of $111In-labeled \ antiTac(Fv)-PE38$ and found that it differed from 125I-labeled antiTac(Fv)-PE38. These studies suggest that 111In-labeled antiTac(Fv)-PE38 can be used to trace the fate of antiTac(Fv)-PE38 in humans.

2/7/2 (Item 2 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
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08785022 BIOSIS NO.: 199395074373

Prolongation of graft survival in primate allograft transplantation by yttrium-90-labeled anti-Tac in conjunction with granulocyte colony-stimulating factor.

AUTHOR: Parenteau Gary L; Dirbas Frederick M; Garmestani Kayhan; Brechbiel Martin W; Bukowski Maria A; Goldman Carolyn K; Clark Richard; Gansow Otto A; Waldmann Thomas A(a)

AUTHOR ADDRESS: (a) Metabolism Branch, National Cancer Inst., Bldg. 10, Room 4N115, National Inst. Health, Bethesda,

JOURNAL: Transplantation (Baltimore) 54 (6):p963-968 1992

ISSN: 0041-1337

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

ABSTRACT: High-affinity IL-2 receptors are expressed by T cells activated in response to foreign histocompatibility antigens but not by normal resting T cells. To exploit this difference in IL-2R expression, anti-Tac, a murine monoclonal antibody specific for the IL-2R-alpha subunit, was used to inhibit organ allograft rejection. To enhance its effector function, anti-Tac was armed by chelation with yttrium-90, a pure beta-emitting radionuclide. Animals received no immunosuppression (n=5, group I, controls), unmodified anti-Tac (n=5, 1 mg/kg, q.o.d., group II), or 90Y-anti-tac (n=5, 1.6 mCi/kg divided into four doses, group III). The animals in group IV (n=4) were treated identically to those in group III with the exception that 5 mu-g/kg/dose of granulocyte colony-stimulating factor was administered intramuscularly on the days when the yttrium-90 was given and on postoperative days 12 through 35 in order to reduce hematopoietic toxicity. Mean graft survival +- S.E.M. for the control group was 8.2+-0.5 days as compared with 13.8+-2.1 days (P lt 0.05) for those monkeys treated with unmodified anti-Tac. Graft survival was further prolonged in animals of group III that received 90Y-anti-Tac, with a mean graft survival of 45.0+-11.8 days; however, three of the five monkeys retained viable grafts within this group but died secondary to bone marrow suppression. In comparison, the monkeys in group IV that were treated with G-CSF in combination with 90Y-anti-Tac had a mean graft survival of 49.2+-2.9 days. In contrast to group III there were no deaths in the group (IV) receiving G-CSF. Furthermore, animals in group IV had a reduced magnitude and shortened duration of irradiation-induced neutropenia when compared with that observed in group III animals that did not receive G-CSF. Thus, treatment with 90Y-anti-tac in conjunction with G-CSF may have

102022586 CA: 102(3)22586x JOURNAL
Expression of interleukin 2 receptors on activated human B cells
AUTHOR(S): Waldmann, Thomas A.; Goldmann, Carolyn K.; Robb, Richard J.;
Depper, Joel M.; Leonard, Warren J.; Sharrow, Susan O.; Bongiovanni,
Kathleen F.; Korsmeyer, Stanley J.; Greene, Warner C.
LOCATION: Metab. Branch, Natl. Cancer Inst., Bethesda, MD, 20205, USA
JOURNAL: J. Exp. Med. DATE: 1984 VOLUME: 160 NUMBER: 5 PAGES: 1450-66
CODEN: JEMEAV ISSN: 0022-1007 LANGUAGE: English

8/3/22 (Item 22 from file: 399)
DIALOG(R)File 399:CA SEARCH(R)
(c) 1999 American Chemical Society. All rts. reserv.

101205721 CA: 101(23)205721m JOURNAL
Pseudomonas exotoxin-anti-TAC. Cell-specific immunotoxin active against
cells expressing the human T-cell growth factor receptor
AUTHOR(S): FitzGerald, David J. P.; Waldmann, Thomas A.; Willingham, Mark
C.; Pastan, Ira
LOCATION: Lab. Mol. Biol., Natl. Cancer Inst., Bethesda, MD, 20205, USA
JOURNAL: J. Clin. Invest. DATE: 1984 VOLUME: 74 NUMBER: 3 PAGES:
966-71 CODEN: JCINAO ISSN: 0021-9738 LANGUAGE: English

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8/3/23 (Item 23 from file: 399)
DIALOG(R)File 399:CA SEARCH(R)

(c) 1999 American Chemical Society. All rts. reserv.

101105569 CA: 101(13)105569z JOURNAL
Phorbol diester induces expression of Tac antigen on human acute T
lymphocytic leukemic cells
AUTHOR(S): Greene, Warner C.; Robb, Richard J.; Depper, Joel M.; Leonard,
Warren J.; Drogula, Cynthia; Svetlik, Penny B.; Wong-Staal, Flossie; Gallo,
Robert C.; Waldmann, Thomas A.
LOCATION: Lab. Tumor Cell Biol., Natl. Cancer Inst., Bethesda, MD, USA
JOURNAL: J. Immunol. DATE: 1984 VOLUME: 133 NUMBER: 2 PAGES: 1042-7
CODEN: JOIMA3 ISSN: 0022-1767 LANGUAGE: English

8/3/24 (Item 24 from file: 399)
DIALOG(R)File 399:CA SEARCH(R)
(c) 1999 American Chemical Society. All rts. reserv.

100083564 CA: 100(11)83564u JOURNAL Hairy cell leukemia: a malignant expansion of B cells which express Tac antigen
AUTHOR(S): Greene, Warner C.; Waldmann, Thomas A.; Cossman, Jeffrey; Hsu, Su Ming; Neckers, Leonard M.; Marshall, Sandra L.; Jensen, Jane P.;
Bakhshi, Ajay; Leonard, Warren J.; et al.
LOCATION: Metab. Branch, Natl. Cancer Inst., Bethesda, MD, 20205, USA JOURNAL: UCLA Symp. Mol. Cell. Biol., New Ser. DATE: 1983 VOLUME: 9
NUMBER: Norm. Neoplast. Hematopoiesis PAGES: 501-11 CODEN: USMBD6
LANGUAGE: English

8/3/25 (Item 25 from file: 399)
DIALOG(R)File 399:CA SEARCH(R)
(c) 1999 American Chemical Society. All rts. reserv.

100004419 CA: 100(1)4419z CONFERENCE PROCEEDING
Monoclonal anti-Tac blocks the action and membrane binding of human
interleukin-2
AUTHOR(S): Depper, Joel M.; Leonard, Warren J.; Smith, Kendall A.;
Waldmann, Thomas A.; Greene, Warner C. !

LOCATION: Metab. Branch, Natl. Cancer Inst., Bethesda, MD, USA JOURNAL: Interleukins, Lymphokines, Cytokines, Proc. Int. Lymphokine Workshop, 3rd EDITOR: Oppenheim, Joost J. (Ed), Cohen, Stanley (Ed), Landy, Maurice (Ed), DATE: 1983 PAGES: 19-26 CODEN: 500PAC LANGUAGE: English MEETING DATE: 820000 PUBLISHER: Academic, New York, N. Y

(Item 26 from file: 399) 8/3/26 DIALOG(R) File 399:CA SEARCH(R) (c) 1999 American Chemical Society. All rts. reserv.

CA: 99(11)86331n **JOURNAL** 99086331

Blockade of the interleukin-2 receptor by anti-Tac antibody: inhibition human lymphocyte activation of human lymphocyte activation

AUTHOR(S): Depper, Joel M.; Leonard, Warren J.; Robb, Richard J.;

Waldmann, Thomas A.; Greene, Warner C.

LOCATION: Metab. Branch, Natl. Cancer Inst., Bethesda, MD, 20205, USA JOURNAL: J. Immunol. DATE: 1983 VOLUME: 131 NUMBER: 2 PAGES: 690-6

CODEN: JOIMA3 ISSN: 0022-1767 LANGUAGE: English

3/K/1 (Item 1 from file: 653)
DIALOG(R)File 653:(c) format only 2000 The Dialog Corp. All rts. reserv.

... specific for tumor cells. With the possible exception of the antigens associated with B-cell **lymphoma** against which anti-idiotype antibodies are directed, no tumor-specific antigens suitable as targets for

... possible exceptions are anti-idiotype antibodies, but any such antibody is specific for the B-lymphoma cells of only one individual, and thus must be separately developed and isolated for each...e.g., cancer cells) in vivo have been developed. Examples of such MAbs are anti-TAC, or other interleukin-2 receptor antibodies; 9.2.27 and NR-ML-05 to a 250 kilodalton human melanoma associated proteoglycan...flow of inert gas.

The chelate is then conjugated to a Fab fragment of monoclonal antibody TFS-2 and a Fab fragment of TFS-4 in separate reaction mixtures. The Fab fragments are generated by papain treatment according to conventional procedures.

A buffered solution of the **antibody** fragment (5 mg/mL, 0.5 mL) is added to the purified sup 188 Re-labeled chelate, followed...

3/K/21 (Item 19 from file: 654)
DIALOG(R)File 654:(c) format only 2000 The Dialog Corp. All rts. reserv.

... AIDS, Meningitis, Arthritis) and therapy of tumors expressing IL-2 receptor such as T-cell **leukemia**.

In fact, the administration of B-B10 on the graft-versus-host reaction as produced... present some other examples in addition to those as hereinabove mentioned. For instance, an anti-Tac antibody is humanized by transplantation of nine amino acid residues on the framework in addition to

...activated human T cell as illustrated in Examples 11 and 14.

The humanized B-B10 antibody of the invention is parenterally administered with a dosage of about 0.05-500 mg for treatment and prevention of diseases caused by graft-versus-host reaction or host-versus-graft reaction...

 \dots AIDS, meningitis, arthritis), and therapy of tumors expressing IL-2 receptor such as T-cell ${\bf leukemia}.$

ADVANTAGEOUS EFFECT OF THE INVENTION

1) Half-life of mouse Mab in blood is about...

3/K/47 (Item 45 from file: 654)
DIALOG(R)File 654:(c) format only 2000 The Dialog Corp. All rts. reserv.

OTHER REFERENCES

... Statton, New York, pp. 283-301 (1986).

Hale et al., "Remission Induction in Non-Hodgkin Lymphoma with

- Reshaped Human Monoclonal Antibody CAMPATH-1H," Lancet, Dec. 17, 1988, pp. 1394-1399.
 - Hieter...virus infection in vivo, "Bio/Technology, 9:26-271 (1991).
 - Uchiyama et al., "A monoclonal **antibody** (anti-**Tac**) reactive with activated and functionally mature human T-cells," J. Immunol., 126:1393-1397 (1981...
 - ...heavy chain (B) (SEQ ID NOS:3 and 4) variable regions of the mouse antiTac antibody (upper lines), compared with the human Eu
 antibody (lower lines), not including signal sequences. The three
 CDR's in each chain are underlined... FIG. 7A through FIG. 7D.
 Fluorocytometry of HUT-102 and Jurkat cells stained with anti-Tac
 antibody or humanized anti-Tac antibody followed
 respectively by fluorescein-conjugated goat anti-mouse Ig antibody
 or goat anti-human Ig antibody, as labeled. In each panel, the dotted curve
 shows...
 - ... then with phycoerythrin-conjugated avidin. (B) Fluorocytometry of HUT-102 cells stained with the indicated **antibody**, then with biotinylated anti-**Tac**, and then with phycoerythrin-conjugated avidin.
 - FIG. 9A and FIG. 9B. Schematic diagram of the...NOS:7 and 8) chain variable regions of the PDL and CDR-only humanized anti-Tac antibodies. The PDL sequence is shown on the upper line, the CDR-only sequence below. Amino...
 - ... light (B) (SEQ ID NO:10) chain variable regions of the CDR-only humanized anti-Tac antibody including signal sequences. Oligonucleotides used for gene synthesis are marked by solid lines: above, for...
 - ... 12. FACS analysis of HUT-102 cells stained with PDL and CDR-only humanized anti-Tac antibodies and negative control antibody Fd79.
 - FIG. 13. Competition by mouse, PDL humanized, and CDR-only humanized anti-Tac antibodies with binding of radioiodinated mouse anti-Tac antibody to HUT-102 cells.
 - FIG. 14. Scheme for anchored polymerase chain reaction (PCR) cloning of ...IL-2 stimulated proliferation of human PHA blasts by humanized mikbeta 1+humanized anti-Tac antibodies. No antibody added (quadrature), 2 ug each of humanized mik- beta 1 and humanized anti-Tac added
 - ...human antibody (e.g., A.T.C.C. Accession No. CRL 9688 secretes an anti-Tac chimeric antibody), although other mammalian species may be

As used herein, the term "humanized" immunoglobulin refers...2 receptor's structure and function is due to the development of specifically reactive monoclonal antibodies. In particular, one mouse monoclonal antibody, known as anti-Tac (Uchiyama, et al., J. Immunol. 126, 1393 (1981)) has been used to show that IL...

...the IL-2 receptor (Herrmann, et al., J. Exp. Med. 162, 1111 (1985)).

The anti-**Tac** monoclonal **antibody** has also been used to define lymphocyte functions that require IL-2 interaction, and has...

... of cytotoxic and suppressor T lymphocytes in cell culture. Also, based on studies with anti-Tac and other antibodies, a variety of disorders are now associated with improper IL-2 receptor expression by T-cells, in particular adult T-cell leukemia.

More recently, the IL-2 receptor has been shown to be an ideal ... approaches to T-cell mediated diseases. It has been proposed that IL-2 receptor specific **antibodies**, such as the anti-Tac monoclonal **antibody**, can be used either alone or as an immunoconjugate (e.g., with Ricin A, isotopes...

... the IL-2 receptor. These agents can, for example, theoretically eliminate IL-2 receptor-expressing **leukemic** cells, certain B-cells, or activated T-cells involved in a disease state, yet allow...

... response by activated T-cells. Indeed, clinical trials have been initiated using, e.g., anti-Tac antibodies (see, generally, Waldmann, T., et al., Cancer Res. 45, 625 (1985), Waldmann, T., Science 232 herein by reference).

Unfortunately, the use of the anti-Tac and other non-human monoclonal antibodies have certain drawbacks, particularly in repeated therapeutic regimens as explained below. Mouse monoclonal antibodies, for...

 \dots and lack other important immunoglobulin functional characteristics when used in humans.

Perhaps more importantly, anti-Tac and other non-human monoclonal antibodies contain substantial stretches of amino acid sequences that will be immunogenic when injected into a...binding to a desired epitope on the human IL-2 receptor, such as the anti-Tac monoclonal antibody. The DNA segments encoding these regions will typically be joined to DNA segments encoding appropriate...selected primarily based on fluid volumes, viscosities, etc., in accordance with the particular mode of administration selected.

Thus, a typical pharmaceutical composition for injection could be made up to contain 1 ml sterile buffered water, and 1 to 50 mg of antibody. A typical composition for intravenous infusion could be made up to contain 250 ml of sterile Ringer's solution, and 150 mg of antibody. Actual methods for preparing parenterally administrable compositions will be known or apparent to those skilled in the art and are described...

... 15th ed., Mack Publishing Company, Easton, Pa. (1980), which is incorporated herein by reference.

The antibodies of this invention can be frozen or lyophilized for storage and reconstituted in a suitable... the disease and its complications. An amount adequate to accomplish this is defined as a "therapeutically effective dose." Amounts effective for this use will depend upon the severity of the infection...

... the patient's own immune system, but generally range from about 1 to about 200 mg of antibody per dose, with dosages of from 5 to 25 mg being more commonly used. It...of health and general level of immunity, but generally range from 0.1 to 25 mg per dose, especially 0.5 to 2.5 mg per dose. A preferred prophylactic use is for the prevention of kidney transplant rejection.

Single or multiple administrations of the compositions can be carried out with dose levels and pattern being selected by the treating physician. In any event, the pharmaceutical formulations should provide a quantity of the antibody(ies) of this invention sufficient to effectively treat the patient.

Human-like **antibodies** of the present invention can further find a wide variety of utilities in vitro. By way of example, the **antibodies** can be utilized for T-cell typing, for isolating specific IL-2 receptor bearing cells... human-like antibody binding to p75 of the IL-2 receptor,

i.e., humanized anti-Tac plus humanized mik- beta 1.

Human-like antibodies of the present invention can further find a wide variety of utilities in vitro. By... cell type, such as cells expressing an HSV epitope.

The compositions containing the present humanized antibodies or a cocktail thereof can be administered for prophylactic and/or therapeutic treatments. In therapeutic application, compositions are administered to a patient already suffering from HSV infection, in an amount sufficient to cure or...

...the disease and its complications. An amount adequate to accomplish this is defined as a "therapeutically effective dose." Amounts effective for this use will depend upon the severity of the infection...

... the patient's own immune system, but generally range from about 1 to about 200 mg of antibody per dose, with dosages of from 5 to 25 mg being more commonly used. It must be kept in mind that the materials of this... humanized immunoglobulins of this invention, it is possible and may be felt desirable by the treating physician to administer substantial excesses of these antibodies.

In prophylactic applications, compositions containing the present immunoglobulins or a cocktail thereof are **administered** to a patient not already in a disease state to enhance the patient's resistance...

...of health and general level of immunity, but generally range from 0.1 to 25 mg per dose. A preferred prophylactic use is for the prevention of herpes in immunocompromised patients, such as organ transplant recipients.

Single or multiple administrations of the compositions can be carried out with dose levels and pattern being selected by the treating physician. In any event, the pharmaceutical formulations should provide a quantity of the antibody(ies) of this invention sufficient to effectively treat the patient.

Humanized antibodies of the present invention can further find a wide variety of utilities in vitro. By way of example, the antibodies can be utilized for detection ...about 10,000-15,000 new cases of myeloid (also called non-lymphocytic or granulocytic) leukemia in the U.S. per year (Cancer Facts & Figures, American Cancer Society, 1987). There are two major forms of myeloid leukemia: acute myelogenous leukemia (AML) and chronic myelogenous leukemia (CML). Despite treatment with chemotherapy, long-term survival in patients with AML is less than...
... Hematology 4, 221 (1986)), and survival with CML and related diseases such as chronic myelomonocytic leukemia (CMML), chronic monocytic leukemia (CMMOL) and myelodysplastic syndrome (MDS) is even lower.

The p67 protein or CD33 antigen is found on the surface of progenitors of myeloid cells and of the **leukemic** cells of most cases of AML, but not on lymphoid cells or non-hematopoietic cells124 (1983) and Griffin et al., **Leukemia** Research 8, 521 (1984), both of which are incorporated herein by reference).

Another antibody that binds to CD33 is M195 (Tanimoto et al., **Leukemia** 3, 339 (1989) and Scheinberg et al., **Leukemia** 3, 440 (1989), both of which are incorporated herein by reference). The reactivity of M195...

... 9, 207 (1990)). M195 radiolabeled with iodine-131 was found to rapidly and specifically target **leukemic** cells in both the blood and bone marrow.

Unfortunately, the use of non-human monoclonal...needs.

The present invention provides novel compositions useful, for example, in

the treatment of myeloid **leukemia** -related human disorders, the compositions containing humanized immunoglobulins specifically capable of binding to CD33 antigen...form, or together with a chemotherapeutic agent such as cytosine arabinoside or daunorubicin active against **leukemia** cells, or complexed with a radionuclide such as iodine-131. All of these compounds will be particularly useful in treating **leukemia** and myeloid cell-mediated disorders. The humanized immunoglobulins or their complexes can be prepared in...

...to 10 sup 10 M sup -1 or stronger, are capable of, e.g., destroying leukemia cells. The humanized immunoglobulins will have a human framework and will have one or ...additional agents (e.g., 6-thioguanine) well-known to those skilled in the art for leukemia treatment may also be utilized (see, Hoffbrund & Pettit., op. cit.).

A preferred pharmaceutical composition of the present invention comprises the use of the subject immunoglobulins in immunotoxins to kill **leukemia** cells. ...examples are offered by way of illustration, not by limitation.

EXPERIMENTAL Example 1

Humanized anti-Tac Antibody

Design of Genes for Humanized anti-Tac Light and Heavy Chains

The sequence of the human antibody Eu (Sequences of Proteins of Immunological Interest, E. Kabat et al., U.S. Dept. of...

- ... humanized antibody, because the amino acid sequence of the heavy chain variable region of anti-Tac is more homologous to the heavy chain of this antibody than to any other complete heavy chain variable region sequence in the National Biomedical Foundation...heavy chain (amino acids 30 and 67); or
- (4) 3-dimensional modeling of the anti-Tac antibody suggested that the amino acid was physically close to the antigen binding region (amino acids...do not express the IL-2 receptor (FIG. 7D). As controls, the original mouse anti-Tac antibody was also used to stain these cells, giving similar results.

For the next experiments, cells...Laboratories, Inc., Richmond, Calif.) according to standard techniques. To determine the affinity of relative to the original anti-Tac humanized antibody antibody, a competitive binding experiment was performed. About 5X10 sup 5 HUT-102 cells were incubated with known quantities (10-40 ng) of the anti-**Tac** antibody and the humanized anti-Tac antibody for 10 min at 4 degree(s) C. Then 100 ng of biotinylated (10-40 ng) of the anti-Tac anti-Tac of increasing amounts antibody as competitor in the first step decreased the amount of biotinylated anti-Tac that could...

...with the biotinylated anti-Tac, thus decreasing fluorescence more.

Example 2

A Second Humanized anti-Tac Antibody

Higher Level Expression of the Humanized anti-Tac Antibody

Three new plasmid vectors were prepared for expression of the humanized antibodies. The plasmid pVgl...Xba I sites of the plasmid vectors pVk and pVgl. To express the humanized anti-Tac antibody, the light chain encoding plasmid was introduced by electroporation into SP2/0 mouse

myeloma cells...

... supernatant of transfected cells by protein A sepharose chromatography.

Construction of the Second Humanized anti-Tac Antibody

To determine whether it was actually necessary to use the mouse anti- Tac amino acids in categories (2)-(4) in the humanized anti-Tac antibody to retain binding affinity, a second humanized anti-Tac antibody was constructed. In the second antibody, only mouse anti-Tac amino acids in Category (1), i.e., in the CDR's themselves, were used, with...

...coming from the human Eu framework. For purposes of this discussion, the original humanized anti-Tac antibody will be called the "PDL humanized antibody," and the second humanized antibody will be called the "CDR-only humanized antibody." The amino...detectably bind the IL-2 receptor.

Binding of the PDL and CDR-only humanized anti-Tac antibodies to the $\mbox{IL-2}$ receptor were also compared in a competitive binding assay. Approximately 4X10 sup 5 HUT-102 cells were incubated with 1.5 ng of radioiodinated mouse anti-Tac antibody (7X10 sup 6 cpm/ug) and varying amounts of each humanized antibody (4 to 512...were collected by centrifugation and radioactivity was measured. The relative binding by the two humanized antibodies and by mouse anti-Tac is shown in a plot of bound/free labelled antibody versus competitor concentration (FIG. 13). The PDL humanized anti-Tac antibody affinity for IL-2 receptor is essentially equal to that of the mouse anti-Tac antibody , because it competes about equally well. But competition by the CDR-only humanized anti-Tac antibody to IL-2 receptor was undetectable at the antibody concentrations used, indicating a binding affinity reduction of at least 100-fold as compared to the PDL humanized anti-Tac antibody. Because the sequences of the PDL and CDR humanized anti-Tac antibodies differ only at positions where mouse framework residues in categories (2)-(4) were used in...of herpes simplex virus (Metcalf et al., Intervirology 29, 39 (1988)), M195 (Tanimoto et al., Leukemia 3, 339 (1989)) which binds to the CD33 antigen, mik- beta 1 (Tusdo et al...of the humanized antibody, because the amino acid sequence of the heavy chain of anti-Tac is more homologous to the heavy chain of this antibody than to any other heavy chain sequence in the National Biomedical Foundation Protein Identification Resourceamino acids 30 and 67).

(4) 3-dimensional modeling of the anti-Tac antibody suggested that the amino acid was physically close to the antigen binding region (amino acids...do not express the IL-2 receptor (FIG. 7D). As controls, the original mouse anti-Tac antibody was also used to stain these cells (FIG. 7B and FIG. 7C), giving similar resultsTo determine the affinity of the humanized antibody relative to the original anti-Tac antibody, a competitive binding experiment was performed. About 5X10 sup 5 HUT-102 cells were incubated with known quantities (10-40 ng) of the anti-Tac antibody and the humanized anti-Tac antibody for 10 min at 4 degree(s) C. Then 100 ng of biotinylated anti-Tac...

...fluorescence on a FACSCAN cytofluorometer.

Use of increasing amounts (10-40 ng) of the anti-Tac antibody as competitor in the first step decreased the amount of biotinylated anti-... if one had greater affinity, it would have more effectively competed with the biotinylated anti-Tac, thus decreasing fluorescence more.

Biological Properties of the Humanized Antibody

For optimal use in treatment of human disease, the humanized antibody should be able to...

...cell such as a macrophage that can lyse the target. To determine whether the humanized antibody and the original mouse anti-Tac antibody can mediate ADCC, a chromium release assay was performed by standard methods. Specifically, human leukemia HUT-102 cells, which express the IL-2 receptor, were incubated with sup 51 Cr...

... this radionuclide. The HUT-102 cells were then incubated with an excess of either anti-Tac or humanized anti-Tac antibody. The HUT-102 cells were next incubated for 4 hrs with either a 30:1...

... is likely to be more efficacious than the original mouse antibody in treating T-cell leukemia or other T-cell mediated diseases.

 $$\operatorname{\texttt{TABLE}}$$ 3Higher Level Expression of the Humanized anti- $\operatorname{\texttt{Tac}}$ Antibody

Two new plasmid vectors were prepared for expression of the humanized antibody. The plasmid pVgl... Xba I sites of the plasmid vectors pVk and pVGl. To express the humanized anti-Tac antibody, the light chain encoding plasmid was introduced by electroporation into SP2/0 mouse myeloma cells...

... immunoglobulins of the present invention offer numerous advantages of other human IL-2 receptor-specific **antibodies**. In comparison to anti-**Tac** mouse monoclonal **antibodies**, the present human-like immunoglobulin can be more economically produced and contain substantially less foreign... antibody was within about 2-fold of the affinity of the mouse mik- beta 1 **antibody**.

The ability of humanized mik- beta 1 plus humanized anti-Tac antibody to inhibit IL-2 stimulated proliferation of human lymphocytes was determined. Human mononuclear cells, collected...of the antibodies will have strong immunosuppressive properties. Humanized mikbeta 1 plus humanized anti-Tac inhibited proliferation much more strongly than did either antibody alone.

From the foregoing, it will be appreciated that the humanized immunoglobulins of the present... ... First and second polynucleotides according to claim 1, wherein the donor immunoglobulin is the anti-Tac antibody or the M195 antibody.

- 5. First and second polynucleotides according to claim 1 wherein the acceptor immunoglobulin heavy and... second polynucleotides according to claim 10 or 11, wherein the donor immunoglobulin is the anti-Tac antibody or the M195 antibody.
- 15. A vector comprising first and second polynucleotides according to claim $10\,.$
 - 16. A vector...

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...for dividing cells.

Chemotherapy for cancer has improved survival in some cancers such as childhood **leukemia** but the survival rate for advanced (metastatic) solid tumors has not dramatically increased in over...cell division. (Cancer, Edited by V. T. Devita et al. 1985 p50 supra).

A monoclonal **antibody Tac** has been reported (Uchiyama et al. J. Immunol. 1981, 126 1393-1397). It appears to...

... interleukin 2 receptor present on activated T lymphocytes but not present on the majority of leukemia cells. However CBL1 has a different antigenic specificity to the Tac antibody. CBL1 reacts with leukemia cell lines but Tac does not. The molecular weight of the CBL1 antigenic determinant is 15,000 daltons. This... antibodies will also find use in detection of malignant cells and in immunological classification of leukemia. Leukemias can be divided into immunological subsets depending on the particular surface antigens displayed by the...

... CBL1 antibody is reactive with three out of the four generally recognised morphological types of leukemia, being nonreactive with chronic lymphocytic leukemia (CLL). CBL1 would thus be useful, in combination with other antibodies having different specificities, in clinical diagnosis of a patient having leukemia. Malignant lymphocyte cells which are incapable of reacting with CBL1 would be presumed to be... old female Balb/c mice (Simonsen Labs) were immunised intravenously with 2X10 sup 6 acute leukemia cells weekly for 3 weeks. Three days after the final injection the spleens were removed...by exclusion gel filtration on S300 (Sephadex sup R). Materials used in characterisation of CBL1.

Leukemia Cells:

Heparinised peripheral blood samples were drawn from children and adults with active leukemia. The acute lymphocytic leukemia (ALL,) end acute myelocytic leukemia (AML) patients had peripheral blood blast counts greater than 90%. Leukemia cells from patients and peripheral blood lymphocytes from healthy donors were isolated by Ficoll-Hypaque sup R density gradient centrifugation. Leukemia cells were stored in liquid nitrogen.

Cell lines were grown in suspension cultures in RPMI...
... of human cell types. The cells that were highly positive were cultured solid tumor cells, leukemia cells, activated lymphocytes and monocytes. Most normal non-dividing tissue cells were not reactive. Therefore...blood monocutes

>95%

15 >80%
Peripheral blood activated
6 >60%
lymphocytes

Malignant cells - Type Acute myeloid leukemias 25

Acute lymphoid **leukemias** 28 >95%

28 >9 Chronic myeloid blast crisis

10 >95% Chronic lymphocytic leukemias

20 <2 Cultured leukemia cell lines,

fultured **leukemia** cell lines, 10 >95%

Reh, CEM, HSB2, Daudi, Raji, HL60, KG, JM, Molt 4, BJABE. Human Studies

Nineteen patients with severe steroid resistant kidney graft rejections were treated with the monoclonal antibody. Five mg in 200 ml of saline ...was given intravenously on 9 successive days. None of the 19 patients developed fever, vomiting, treatment created infections or other side effects. All patients had previously been immunosuppressed with conventional antirejection...abdomen causing a large tumor abscess on the left groin and protruding rectal tumor was treated with CBL1 purified monoclonal antibody. Following ten daily treatments of 20 mg per day the tumor abscess healed completely and the protruding rectal tumor receded.

3/K/54 (Item 52 from file: 654)
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... therapy is further discussed in the two articles; Kozak et al, "Bismuth-212-labeled anti-**Tac** monoclonal **antibody** : Alpha-particle-emitting Radionuclides ... diseases, cardiovascular diseases, inflammatory conditions and other pathological conditions.

The cancer states include carcinomas, sarcomas, leukemias, lymphomas, myelomas and neural tumors.

The infectious diseases include those caused by invading microbes or parasites...Respiratory syncytial virus Varicella-Zoster virus Hepatitis B virus Measles virus Adenovirus Human T-cell leukemia viruses Epstein-Barr virus Murine leukemia virus* Mumps virus Vesicular stomatitis virus Sindbis virus Lymphocytic choriomeningitis virus Wart virus Blue tongue virus Sendai virus Feline leukemia virus* Reo virus Polio virus Simian virus 40* Mouse mammary tumor virus* Dengue virus Rubella...

... antibody fragments reactive with a tumor associated antigens present on carcinoma or sarcoma cells or lymphomas. Such antibodies are disclosed, e.g., in Goldenberg et al., Journal of Clinical Oncology, Vol... in vivo with a wide variety of other radiometals of use in cancer detection and therapy. Thus, a patient could be imaged with the In-111 antibody conjugate of the ligand and thereafter treated with the yttrium complex of the same antibody chelate conjugate, thus facilitating calculation of the dose of radioactivity transported to the patient's tumor and so increasing likelihood of the effective application of the therapy.

The metal chelate conjugated **antibodies** of this invention can be **administered** in vivo in any suitable pharmaceutical carrier. As noted earlier, a physiologic normal saline solution...

...include a minor amount of carrier protein such as human serum albumin to stabilize the **antibody**. The concentration of metal chelate ... conjugated antibodies within the solution will be a matter of choice. Levels of 0.5 **mg** per ml are readily attainable but the concentrations may vary considerably depending upon the specifics...

... addition to these located at the site, the circulating antigens can be removed prior to **treatment**. Such removal of antigens can be accomplished, of example, by the use of unlabeled **antibodies**, or by plasmapheresis in which the patient's serum is **treated** to removed antigens.

A physiological solution of the protein conjugate is advantageously

- metered into sterile metal free vials, e.g., at a unit dosage of about 0.1--100 mg protein conjugate/vial, and the vials are either stoppered, sealed and stored at low temperature...
- ... claim 1, wherein said antibody or antibody fragment specifically binds to an antigen associated with lymphomas, carcinomas, sarcomas, leukemias, myelomas or neural tumors.
 - 4. The conjugate of claim 1, wherein said marker is a...
- ... antibody or F(ab') sub 2 antibody fragment specifically binds to an antigen associated with **lymphomas**, carcinomas, sarcomas, **leukemias**, myelomas or neural tumors.
 - 16. A method for preparing a precursor conjugate of a therapeutic...
- 3/K/62 (Item 60 from file: 654)
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OTHER REFERENCES

...a mouse", Nature 321:522-525 (1986).

Hale et al., "Remission Induction in Non-Hodgkin Lymphoma with Reshaped Human Monoclonal Antibody CAMPATH-1H", Lancet Dec. 17, 1988, pp. 1394-1399.

Chothia...

- ...heavy chain (B) [SEQ ID NOS:3 and 4] variable regions of the mouse anti-Tac antibody (upper lines), compared with the human Eu antibody (lower lines), not including signal sequences. The three CDR's in each chain are underlined...
- ... FIG. 7A through FIG. 7D. Fluorocytometry of HUT-102 and Jurkat cells stained with anti-Tac antibody or humanized anti-Tac antibody followed respectively by fluorescein-conjugated goat anti-mouse Ig antibody or goat anti-human Ig antibody, as labeled. In each panel, the dotted curve shows...
- ... then with phycoerythrin-conjugated avidin. (B) Fluorocytometry of HUT-102 cells stained with the indicated **antibody**, then with biotinylated anti-**Tac**, and then with phycoerythrin-conjugated avidin.
- FIG. ... NOS:7 and 8] chain variable regions of the PDL and CDR-only humanized anti-Tac antibodies. The PDL sequence is shown on the upper line, the CDR-only sequence below. Amino...
- ... light (B) [SEQ ID NO:10] chain variable regions of the CDR-only humanized anti-Tac antibody including signal sequences. Oligonucleotides used for gene synthesis are marked by solid lines: above, for...
- ... 12. FACS analysis of HUT-102 cells stained with PDL and CDR-only humanized anti-Tac antibodies and negative control antibody Fd79.
- FIG. 13. Competition by mouse, PDL humanized, and CDR-only humanized anti-Tac antibodies with binding of radioiodinated mouse anti-Tac antibody to HUT-102 cells.
- FIG. 14. Scheme for anchored polymerase chain reaction (PCR) cloning of ...IL-2 stimulated proliferation of human PHA blasts by humanized mik- beta 1+humanized anti-Tac antibodies. No antibody added (quadrature), 2 ug each of humanized mik- beta 1 and humanized anti-Tac added...

...human antibody (e.g., A.T.C.C. Accession No. CRL 9688 secretes an anti-Tac chimeric antibody), although other mammalian species may be

As used herein, the term "humanized" immunoglobulin refers...2 receptor's structure and function is due to the development of specifically reactive monoclonal antibodies. In particular, one mouse monoclonal antibody, known as anti-Tac (Uchiyama, et al., J. Immunol. 126, 1393 (1981)) has been used to show that IL...

...the IL-2 receptor (Herrmann, et al., J. Exp. Med. 162, 1111 (1985)).

The anti-Tac monoclonal antibody has also been used to define lymphocyte functions that require IL-2 interaction, and has...

... of cytotoxic and suppressor T lymphocytes in cell culture. Also, based on studies with anti-Tac and other antibodies, a variety of disorders are now associated with improper IL-2 receptor expression by T-cells, in particular adult T-cell leukemia.

More recently, the IL-2 receptor has been shown to be an ideal target for

- ... approaches to T-cell mediated diseases. It has been proposed that IL-2 receptor specific **antibodies**, such as the anti-Tac monoclonal **antibody**, can be used either alone or as an immunoconjugate (e.g., with Riçin A, isotopes...
- ... the IL-2 receptor. These agents can, for example, theoretically eliminate IL-2 receptor-expressing **leukemic** cells, certain B-cells, or activated T-cells involved in a disease state, yet allow...
- ... response by activated T-cells. Indeed, clinical trials have been initiated using, e.g., anti-Tac antibodies (see, generally, Waldmann, T., et al., Cancer Res. 45, 625 (1985), Waldmann, T., Science 232
- ...1989), all of which are incorporated herein by reference).

Unfortunately, the use of the anti-Tac and other non-human monoclonal antibodies have certain drawbacks, particularly in repeated therapeutic regimens as explained below. Mouse monoclonal antibodies, for...

 \dots and lack other important immunoglobulin functional characteristics when used in humans.

Perhaps more importantly, anti-Tac and other non-human monoclonal antibodies contain substantial stretches of amino acid sequences that will be immunogenic when injected into a...binding to a desired epitope on the human IL-2 receptor, such as the anti-Tac monoclonal antibody. The DNA segments encoding these regions will typically be joined to DNA segments encoding appropriate...selected primarily based on fluid volumes, viscosities, etc., in accordance with the particular mode of administration selected.

Thus, a typical pharmaceutical composition for injection could be made up to contain 1 ml sterile buffered water, and 1 to 50 mg of antibody. A typical composition for intravenous infusion could be made up to contain 250 ml of sterile Ringer's solution, and 150 mg of antibody. Actual methods for preparing parenterally administrable compositions will be known or apparent to those skilled in the art and are described...

... 15th ed., Mack Publishing Company, Easton, Pa. (1980), which is incorporated herein by reference.

The antibodies of this invention can be frozen or lyophilized for

storage and reconstituted in a suitable...

...the disease and its complications. An amount adequate to accomplish this is defined as a "therapeutically effective dose." Amounts effective for this use will depend upon the severity of the infection...

... the patient's own immune system, but generally range from about 1 to about 200 mg of antibody per dose, with dosages of from 5 to 25 mg being more commonly used. It...

...of health and general level of immunity, but generally range from 0.1 to 25 mg per dose, especially 0.5 to 2.5 mg per dose. A preferred prophylactic use is for the prevention of kidney transplant rejection.

Single or multiple administrations of the compositions can be carried out with dose levels and pattern being selected by the treating physician. In any event, the pharmaceutical formulations should provide a quantity of the antibody(ies) of this invention sufficient to effectively treat the patient.

Human-like antibodies of the present invention can further find a wide variety of utilities in vitro. By way of example, the antibodies can be utilized for T-cell typing, for isolating specific IL-2 receptor bearing cells... human-like antibody binding to p75 of the IL-2 receptor, i.e., humanized anti-Tac plus humanized mik- beta 1.

Human-like **antibodies** of the present invention can further find a wide variety of utilities in vitro. By... cell type, such as cells expressing an HSV epitope.

The compositions containing the present humanized antibodies or a cocktail thereof can be administered for prophylactic and/or therapeutic treatments. In therapeutic application, compositions are administered to a patient already suffering from HSV infection, in an amount sufficient to cure or...

...the disease and its complications. An amount adequate to accomplish this is defined as a "therapeutically effective dose." Amounts effective for this use will depend upon the severity of the infection...

... the patient's own immune system, but generally range from about 1 to about 200 mg of antibody per dose, with dosages of from 5 to 25 mg being more commonly used. It must be kept in mind that the materials of this... humanized immunoglobulins of this invention, it is possible and may be felt desirable by the treating physician to administer substantial excesses of these antibodies.

In prophylactic applications, compositions containing the present immunoglobulins or a cocktail thereof are **administered** to a patient not already in a disease state to enhance the patient's resistance...

...of health and general level of immunity, but generally range from 0.1 to 25 mg per dose. A preferred prophylactic use is for the prevention of herpes in immunocompromised patients, such as organ transplant recipients.

Single or multiple administrations of the compositions can be carried out with dose levels and pattern being selected by the treating physician. In any event, the pharmaceutical formulations should provide a quantity of the antibody(ies) of this invention sufficient to effectively treat the patient.

Humanized **antibodies** of the present invention can further find a wide variety of utilities in vitro. By way of example, the **antibodies** can be utilized for detection of HSV antigens, for isolating specific HSV infected cells or...

 \dots about 10,000-15,000 new cases of myeloid (also called non-lymphocytic or

granulocytic) leukemia in the U.S. per year (Cancer Facts & Figures, American Cancer Society, 1987). There are two major forms of myeloid leukemia: acute myelogenous leukemia (AML) and chronic myelogenous leukemia (CML). Despite treatment with chemotherapy, long-term survival in patients with AML is less than...

... Hematology 4, 221 (1986)), and survival with CML and related diseases such as chronic myelomonocytic **leukemia** (CMML), chronic monocytic **leukemia** (CMMOL) and myelodysplastic syndrome (MDS) is even lower.

The p67 protein or CD33 antigen is found on the surface of progenitors of myeloid cells and of the **leukemic** cells of most cases of AML, but not on lymphoid cells or non-hematopoietic cells...

... L4B3, L1B2 and MY9 (Andrews et al., Blood 62, 124 (1983) and Griffin et al., **Leukemia** Research 8, 521 (1984), both of which are incorporated herein by reference).

Another antibody that binds to CD33 is M195 (Tanimoto et al., Leukemia 3, 339 (1989) and Scheinberg et al., Leukemia 3, 440 (1989), both of which are incorporated herein by reference). The reactivity of M195...

... 9, 207 (1990)). M195 radiolabeled with iodine-131 was found to rapidly and specifically target **leukemic** cells in both the blood and bone marrow.

Unfortunately, the use of non-human monoclonal...

...needs.

The present invention provides novel compositions useful, for example, in the treatment of myeloid leukemia -related human disorders, the compositions containing humanized immunoglobulins specifically capable of ... form, or together with a chemotherapeutic agent such as cytosine arabinoside or daunorubicin active against leukemia cells, or complexed with a radionuclide such as iodine-131. All of these compounds will be particularly useful in treating leukemia and myeloid cell-mediated disorders. The humanized immunoglobulins or their complexes can be prepared in...

- ...to 10 sup 10 M sup -1 or stronger, are capable of, e.g., destroying **leukemia** cells. The humanized immunoglobulins will have a human framework and will have one or more...
- ... additional agents (e.g., 6-thioguanine) well-known to those skilled in the art for **leukemia** treatment may also be utilized (...of the present invention comprises the use of the subject immunoglobulins in immunotoxins to kill **leukemia** cells. Immunotoxins are characterized by two components and are particularly useful for killing selected cells... examples are offered by way of illustration, not by limitation.

EXPERIMENTAL

EXAMPLE 1

Humanized anti-Tac antibody

Design of genes for humanized anti-Tac light and heavy chains

The sequence of the human **antibody** Eu (Sequences of Proteins of Immunological Interest, E. Kabat et al., U.S. Dept. of...

... humanized antibody, because the amino acid sequence of the heavy chain variable region of anti-Tac is more homologous to the heavy chain of this antibody than to any other complete heavy chain variable region

sequence in the National Biomedical Foundation...heavy chain (amino acids 30 and 67); or

(4) 3-dimensional modeling of the anti-Tac antibody suggested that the amino acid was physically close to the antigen binding region (amino acids...do not express the IL-2 receptor (FIG. 7D). As controls, the original mouse anti-Tac antibody was also used to stain these cells, giving similar results.

For the next experiments, cells...

... Laboratories, Inc., Richmond, Calif.) according to standard techniques. To determine the affinity of the humanized **antibody** relative to the original anti-**Tac antibody**, a competitive binding experiment was performed. About 5X10 sup 5 HUT-102 cells were incubated with known quantities (10-40 ng) of the anti-**Tac antibody** and the humanized anti-**Tac antibody** for 10 min at 4 degree(s) C. Then 100 ng of biotinylated anti-Tac...

...fluorescence on a FACSCAN cytofluorometer.

Use of increasing amounts (10-40 ng) of the anti-Tac antibody as competitor in the first step decreased the amount of biotinylated anti-Tac that could... with the biotinylated anti-Tac, thus decreasing fluorescence more.

EXAMPLE 2

A second humanized anti-Tac antibody

Higher level expression of the humanized anti-Tac antibody

Three new plasmid vectors were prepared for expression of the humanized antibodies. The plasmid pVg1...

... Xba I sites of the plasmid vectors pVk and pVgl. To express the humanized anti-Tac antibody, the light chain encoding plasmid was introduced by electroporation into SP2/0 mouse myeloma cells...

... supernatant of transfected cells by protein A sepharose chromatography.

Construction of the second humanized anti-Tac antibody

To determine whether it was actually necessary to use the mouse anti-Tac amino acids in categories (2)-(4) in the humanized anti-Tac antibody to retain binding affinity, a second humanized anti-Tac antibody was constructed. In the second antibody, only mouse anti-Tac amino acids in Category (1), i.e., in the CDR's themselves, were used, with...

...coming from the human Eu framework. For purposes of this discussion, the original humanized anti-Tac antibody will be called the "PDL humanized antibody," and the second humanized antibody will be called the "CDR-only humanized antibody." The amino...detectably bind the IL-2 receptor.

Binding of the PDL and CDR-only humanized anti-**Tac antibodies** to the IL-2 receptor were also compared in a competitive binding assay. Approximately 4X10 sup 5 HUT-102 cells were incubated with 1.5 ng of radioiodinated mouse anti-**Tac antibody** (7X10 sup 6 cpm/ug) and varying amounts of each humanized antibody (4 to 512...

... were collected by centrifugation and radioactivity was measured. The relative binding by the two humanized **antibodies** and by mouse anti-**Tac** is shown in a plot of bound/free labelled **antibody** versus competitor concentration (FIG. 13). The PDL humanized anti-**Tac antibody** affinity for IL-2 receptor is essentially equal to that of

the mouse anti-Tac antibody, because it competes about equally well. But competition by the CDR-only humanized anti-Tac antibody to IL-2 receptor was undetectable at the antibody concentrations used, indicating a binding affinity reduction of at least 100-fold as compared to the PDL humanized anti-Tac antibody. Because the sequences of the PDL and CDR humanized anti-Tac antibodies differ only at positions where mouse framework residues in categories (2)-(4) were used in...

... of herpes simplex virus (Metcalf et al., Intervirology 29, 39 (1988)), M195 (Tanimoto et al., Leukemia 3, 339 (1989)) which binds to the CD33 antigen, mik- beta 1 (Tusdo et al...of the humanized antibody, because the amino acid sequence of the heavy chain of anti-Tac is more homologous to the heavy chain of this antibody than to any other heavy chain sequence in the National Biomedical Foundation Protein Identification Resource...

- ... Tac heavy chain (amino acids 30 and 67).
- (4) 3-dimensional modeling of the anti-Tac antibody suggested that the amino acid was physically close to the antigen binding region (amino acids...do not express the IL-2 receptor (FIG. 7D). As controls, the original mouse anti-Tac antibody was also used to stain these cells (FIG. 7B and FIG. 7C), giving similar results...
- ... Laboratories, Inc., Richmond, Calif.) according to standard techniques. To determine the affinity of the humanized **antibody** relative to the original anti-**Tac antibody**, a competitive binding experiment was performed. About 5X10 sup 5 HUT-102 cells were incubated with known quantities (10-40 ng) of the anti-**Tac antibody** and the humanized anti-**Tac antibody** for 10 min at 4 degree(s) C. Then 100 ng of biotinylated anti-Tac...
- ...fluorescence on a FACSCAN cytofluorometer.

Use of increasing amounts (10-40 ng) of the anti-Tac antibody as competitor in the first step decreased the amount of biotinylated anti-Tac that could...

... if one had greater affinity, it would have more effectively competed with the biotinylated anti-Tac, thus decreasing fluorescence more.

Biological properties of the humanized antibody

For optimal use in treatment of human disease, the humanized antibody should be able to...

...cell such as a macrophage that can lyse the target. To determine whether the humanized **antibody** and the original mouse anti-**Tac antibody** can mediate ADCC, a chromium release assay was performed by standard methods. Specifically, human **leukemia** HUT-102 cells, which express the IL-2 receptor, were incubated with sup 51 Cr...

... this radionuclide. The HUT-102 cells were then incubated with an excess of either anti-Tac or humanized anti-Tac antibody. The HUT-102 cells were next incubated for 4 hrs with either a 30:1...is likely to be more efficacious than the original mouse antibody in treating T-cell leukemia or other T-cell mediated diseases.

TABLE 3Antibody

Anti-**Tac** 4% <1%

Humanized anti-Tac 24% 23 Higher level expression of the humanized anti-Tac antibody

Two new plasmid vectors were prepared for expression of the humanized antibody. The plasmid pVg1...

 \dots Xba I sites of the plasmid vectors pVk and pVG1. To express the humanized anti-Tac antibody, the light chain encoding plasmid

was introduced by electroporation into SP2/0 mouse myeloma cells...

... immunoglobulins of the present invention offer numerous advantages of other human IL-2 receptor-specific **antibodies**. In comparison to anti-**Tac** mouse monoclonal **antibodies**, the present human-like immunoglobulin can be more economically produced and contain substantially less foreign... antibody was within about 2-fold of the affinity of the mouse mik- beta 1 **antibody**.

The ability of humanized mik- beta 1 plus humanized anti-Tac antibody to inhibit IL-2 stimulated proliferation of human lymphocytes was determined. Human mononuclear cells, collected...

... of the antibodies will have strong immunosuppressive properties. Humanized mik- beta 1 plus humanized anti-Tac inhibited proliferation much more strongly than did either antibody alone.

From the foregoing, it will be appreciated that the humanized immunoglobulins of the present...

- ... 6. A humanized immunoglobulin according to claim 1, wherein said donor immunoglobulin is the anti-Tac antibody.
- 7. A humanized immunoglobulin according to claim 1, wherein said acceptor immunoglobulin heavy and light...

3/K/68 (Item 66 from file: 654)
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OTHER REFERENCES

... et al., "Metabolizable sup 111 In Chelate conjugated anti-iodiotype monoclonal antibody for radioimmunodetection of **lymphoma** in mice," J. Nucl. Med. 12:455-460, 1986.

March, Advanced Organic Chemistry, Second edition...

... including monoclonal antibodies specific for tumor-associated antigens in humans. Among the many such monoclonal **antibodies** that may be used are anti-TAC, or other interleukin-2 receptor **antibodies**; 9.2.27 and NR-ML-05, reactive with the 250 kilodalton human melanoma-associated...0 degree(s) C. ice bath for 2 minutes.

The Fab fragment of a monoclonal **antibody** (10 **mg** in 0.5 mL of PBS) was generated by **treating** the monoclonal **antibody** with papain according to conventional techniques. The monoclonal **antibody**, designated NR-LU-10, recognizes a pancarcinoma antigen. Other proteins may be substituted for the...

3/K/70 (Item 68 from file: 654)
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OTHER REFERENCES

... et al., "Metabolizable sup 111 In chelate conjugated anti-idiotype monoclonal antibody for radioimmunodetection of **lymphoma** in mice", J. Nucl. Med. 12:455-460, 1986.

Meares et al., "Chelate Radiochemistry: cleavable...

...monoclonal anti-bodies specific for tumor-associated antigens in humans. Among the many such monoclonal **antibodies** that may be used are anti-**TAC**, or other interleukin-2 receptor **antibodies**; 9. ...a 0 degree(s) ice bath for 2 minutes.

The Fab fragment of a monoclonal **antibody** (10 **mg** in 0.5 mL of PBS) was generated by **treating** the monoclonal **antibody** with papain according to conventional techniques. The monoclonal **antibody**, designated NR-LU-10, recognizes a pancarcinoma antigen. Other proteins may be substituted for the...

3/K/71 (Item 69 from file: 654)
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...81 [1987], 73).

Uchiyama et al (J. Immunol. 126[1981], 1398) described the first monoclonal **antibody** to recognize the **Tac**-antigen (epitope), the short chain of the interleukin-2 receptor.

European Application 0 241 811...

... activated T and B cells but are inactive against quiescent lymphocytes. One of these monoclonal **antibodies**, an anti-**Tac** analogue, enters into interaction with the **Tac**-antigen, the shorter chain of the receptor molecule. The other monoclonal antibody also obviously enters ...

... has not yet been demonstrated.

European Application 0 240 344 concerns anti-CD 4 monoclonal antibodies and anti-Tac analogues (CD 25). Its activity with respect to immunosuppression and specifically with respect to prevention... monoclonal antibody binds to an epitope of the human interleukin 2 receptor that is not Tac. The antibody is accordingly appropriate for the treatment of and prophylaxis against such diseases as hyperimmune syndrome...

... encephalomyelitis, arthritis, etc.), and for interleukin-2 receptor expressing tumor diseases such as T-cell **leukemia**.

The monoclonal antibody can be employed either as such, coupled with magnetic beads, radioactive substances...
... PBS) with 10 mu l of test residue or control monoclonal antibody. The control monoclonal antibodies in this and in the following examples were the anti-Tac-analogous monoclonal antibodies BF 2 and BG 8 (CRTS, Besancon). The bound monoclonal antibody was detected with an... cells was tested on days 0 and 4 as in Example 5a. The control monoclonal antibodies were the Tac analogues
B-F2: CRTS, Besancon (France)

B-F2: CRTS, Besancon (France) 33B3.1: Inserm U 110, Marseille (France) Clonab IL...risk to infections.

In conjunction with the following clinical-emergency indications, 0.1 to 20 mg and preferably 2.5 to 5 mg per day of the monoclonal antibody B.B.10 were administered to the patients within 30 minutes in the form of an intravenous infusion. The dose...

...selected to attain a plasma level of 0.5 to 5 mu g/ml.

The **treatment** was continued for 3 to 30 days and preferably for 7 to 10 days until the interleukin-2-dependent symptoms subsided. **Treatment** beyond that point surprisingly turned out to be unnecessary.

EXAMPLE 8

The amounts of the monoclonal **antibody** B.B.10 (diluted in human albumin) listed in Table 5 were **administered** at the times cited in the table to a group of 4 patients 4, 11...marrow transplantation on the third one. Due to their age, the dosage of the monoclonal **antibody**

 ${\tt B.B.10}$ was only 2.5 ${\tt mg}$ per day.

Table 6 indicates the doses and duration of treatment.

TABLE 6... ? t s3/3/1,21,47,50,54,62,68,70,71

1/3/1 DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv. 0234528 DBA Accession No.: 99-04629 PATENT Killing cells with immunotoxin - cell lysis using a Pseudomonas sp. exotoxin and single chain antibody scFv fragment fusion, antibody engineering for autoimmune disease and cancer therapy AUTHOR: Fitzgerald D J; Chaudhary V K; Pastan I H; Waldmann T A; Queen C L CORPORATE SOURCE: Washington, DC, USA. PATENT ASSIGNEE: U.S.Dep.Health-Hum.Serv. 1999 PATENT NUMBER: US 5863745 PATENT DATE: 990126 WPI ACCESSION NO.: 99-131300 (9911) PRIORITY APPLIC. NO.: US 461825 APPLIC. DATE: 950605 NATIONAL APPLIC. NO.: US 461825 APPLIC. DATE: 950605 LANGUAGE: English DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv. 0220988 DBA Accession No.: 98-02585 PATENT Fusion protein of antibody Fv fragment and Pseudomonas exotoxin fragment an immunotoxin for use as e.g. an antitumor or immunosuppressive agent AUTHOR: FitzGerald D; Chaudhary V K; Pastan I H; Waldmann T A; Oueen C L CORPORATE SOURCE: Washington, DC, USA. PATENT ASSIGNEE: U.S.Dep.Health-Hum.Serv. 1997 PATENT NUMBER: US 5696237 PATENT DATE: 971209 WPI ACCESSION NO.: 98-041352 (9804) PRIORITY APPLIC. NO.: US 463163 APPLIC. DATE: 950605 NATIONAL APPLIC. NO.: US 463163 APPLIC. DATE: 950605 LANGUAGE: English 1/3/3 DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv. 0164255 DBA Accession No.: 94-06806 Genetically engineered monoclonal antibodies armed with radionuclides radionuclide-carrying anti-Tac monoclonal antibody with reduced immunogenicity (humanized antibody engineering) for interleukin-2 receptor-directed therapy (conference paper) AUTHOR: Waldmann T A CORPORATE AFFILIATE: Nat.Cancer-Inst.Bethesda Nat.Inst.Health-Bethesda CORPORATE SOURCE: Metabolism Branch, Bldg 10, Rm 4N115, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892, USA. JOURNAL: BIOTECH'92 (7, 205-12) 1993 CODEN: 9999Q LANGUAGE: English 1/3/4 DIALOG(R) File 357: Derwent Biotechnology Abs

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0123619 DBA Accession No.: 91-11261 Monoclonal antibodies in diagnosis and therapy - monoclonal antibody production; a review AUTHOR: Waldmann T A CORPORATE SOURCE: Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892, USA. JOURNAL: Science (252, 5013, 1657-62) 1991 CODEN: SCIEAS LANGUAGE: English 1/3/5 DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv. 0110818 DBA Accession No.: 90-13509 IL2-PE664Glu, a new chimeric protein cytotoxic to human-activated T-lymphocytes - human interleukin-2 gene fusion with Pseudomonas exotoxin mutant gene; potential use of chimeric toxin as cytostatic and immunosuppressive AUTHOR: Loberboum-Galski H; Garsia R J; Gately M; Brown P S; Clark R E; Waldmann T A CORPORATE AFFILIATE: Roche CORPORATE SOURCE: Laboratory of Molecular Biology, Division of Cancer Biology and Diagnosis, National Cancer Institute, National Institutes of Health, Bethesda, Maryland 20892, USA. JOURNAL: J.Biol.Chem. (265, 27, 16311-17) 1990 CODEN: JBCHA3 LANGUAGE: English 1/3/6 DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv. 0090949 DBA Accession No.: 89-08940 A recombinant immunotoxin consisting of two antibody variable domains fused to Pseudomonas exotoxin - monoclonal antibody anti-Tac to the p55 subunit of human interleukin-2 receptor AUTHOR: Chaudhary V K; Queen C; Junghans R P; Waldmann T A; FitzGerald D J; Pastan I CORPORATE AFFILIATE: Protein-Design-Labs CORPORATE SOURCE: Laboratory of Molecular Biology, DCBD, National Cancer Institute, National Institutes of Health, Bethesda, Maryland 20892, JOURNAL: Nature (339, 6223, 394-97) 1989 CODEN: NATUAS LANGUAGE: English DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv. 0054663 DBA Accession No.: 86-12511 Radionuclide-conjugated monoclonal antibodies: a synthesis of immunology, inorganic chemistry and nuclear science - development and application in tumor diagnosis etc. AUTHOR: Kozak R W; Waldmann T A; Atcher R W; Gansow O A CORPORATE SOURCE: Metabolism Branch, National Cancer Institute of NIH, Bethesda, MD 20892, USA. JOURNAL: Trends Biotechnol. (4, 10, 259-64) 1986

CODEN: 8921M LANGUAGE: English 1/3/8
DIALOG(R)File 357:Derwent Biotechnology Abs
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0049478 DBA Accession No.: 86-07326

The structure, function, and expression of interleukin-2 receptors on normal and malignant lymphocytes - investigated using a monoclonal antibody

AUTHOR: Waldmann T A

CORPORATE SOURCE: Metabolism Branch, National Cancer Institute, National

Institutes of Health, Bethesda, MD 20892, USA.

JOURNAL: Science (232, 4751, 727-32) 1986

CODEN: SCIEAS LANGUAGE: English

1/3/9

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0005909 DBA Accession No.: 82-04909

A monoclonal antibody that appears to recognize the receptor for human T-cell growth factor; partial characterization of the receptor - purification of the interleukin-2 surface receptor by immunoprecipitation

AUTHOR: Leonard W J; Depper J M; Uchiyama T; Smith K A; Waldmann T A; Greene W C

CORPORATE SOURCE: The Metabolism Branch, National Cancer Institute, NIH, Bethesda, Maryland 20205, USA.

JOURNAL: Nature (300, 5889, 267-69) 1982

CODEN: NATUAS LANGUAGE: English

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              49 AU=WALDMANN THOMAS A
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DIALOG(R) File 5:Biosis Previews(R)
(c) 2000 BIOSIS. All rts. reserv.
          BIOSIS NO.: 200000043153
Patients with chemotherapy-refractory hairy cell leukemia achieve major
  responses to recombinant immunotoxins containing truncated pseudomonas
  exotoxin and targeting either CD25 or CD22.
AUTHOR: Kreitman Robert J(a); Wilson Wyndham H; Margulies Inger(a);
  Stetler-Stevenson Maryalice; Raggio Miranda; FitzGerald David J(a);
  Waldmann Thomas A; Pastan Ira(a
AUTHOR ADDRESS: (a)Laboratory of Molecular Biology, National Cancer
  Institute, NIH, Bethesda, MD**USA
JOURNAL: Blood 94 (10 SUPPL. 1 PART 1):p96a Nov. 15, 1999
CONFERENCE/MEETING: Forty-first Annual Meeting of the American Society of
Hematology New Orleans, Louisiana, USA December 3-7, 1999
SPONSOR: The American Society of Hematology
ISSN: 0006-4971
RECORD TYPE: Citation
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LANGUAGE: English (Item 2 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. 12266169 BIOSIS NO.: 200000019671 Responses in refractory hairy cell leukemia to a recombinant immunotoxin. AUTHOR: Kreitman Robert J; Wilson Wyndham H; Robbins David; Margulies Inger ; Stetler-Stevenson Maryalice; Waldmann Thomas A; Pastan Ira(a AUTHOR ADDRESS: (a) National Cancer Institute, National Institutes of Health, 37 Convent Dr, 37/4E16, MSC 4255, Bethesda, MD, 20892**USA JOURNAL: Blood 94 (10):p3340-3348 Nov. 15, 1999 ISSN: 0006-4971 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English SUMMARY LANGUAGE: English 3/3/3 (Item 3 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. BIOSIS NO.: 199900417059 12122210 Methods to avoid adverse effect of circulating antigen on biodistribution of 125I-labeled antiTac dsFv: Preinjection of intact antibody versus clearance of antigen with avidin-biotin system. AUTHOR: Kobayashi Hisataka; Sun Bao-Fu; Yoo Tae M; Le Nhat; Kim Meyoung-Kon ; Paik Chang H; Pastan Ira; Waldmann Thomas A; Carrasquillo Jorge A (a AUTHOR ADDRESS: (a) Nuclear Medicine Department, National Institutes of Health, 10 Center Dr., Building 10, Rm. 1C-496, Bethesda, MD, 20892-1180 **USA JOURNAL: Journal of Nuclear Medicine 40 (8):p1381-1391 Aug., 1999 ISSN: 0161-5505 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English SUMMARY LANGUAGE: English (Item 4 from file: 5) DIALOG(R) File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. BIOSIS NO.: 199900270961 Favorable effects of glycolate conjugation on the biodistribution of humanized antiTac Fab fragment. AUTHOR: Kobayashi Hisataka; Kim In-sook; Drumm Debbie; Kim Meyoung-kon; Paik David S; Le Nhat; Waldmann Thomas A; Carrasquillo Jorge A(a); Paik Chang H AUTHOR ADDRESS: (a) Department of Nuclear Medicine, National Institutes of Health, 10 Center Dr., MSC 1180, Bldg. 10**USA

Health, 10 Center Dr., MSC 1180, Bldg. 10**USA

JOURNAL: Journal of Nuclear Medicine 40 (5):p837-845 May, 1999

ISSN: 0161-5505

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

SUMMARY LANGUAGE: English

3/3/5 (Item 5 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2000 BIOSIS. All rts. reserv.

11848859 BIOSIS NO.: 199900094968 The pharmacokinetic characteristics of glycolated humanized anti-Tac Fabs are determined by their isoelectric points. AUTHOR: Kobayashi Hisataka; Le Nhat; Kim In-Sook; Kim Meyoung-Kon; Pie Jae-Eun; Drumm Debra; Paik David S; Waldmann Thomas A; Paik Chang H ; Carrasquillo Jorge A(a AUTHOR ADDRESS: (a) NIH, Building 10, Room IC496, 10 Center Dr., MSC 1180, Bethesda, MD 20892-1180**USA JOURNAL: Cancer Research 59 (2):p422-430 Jan. 15, 1999 ISSN: 0008-5472 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English (Item 6 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. BIOSIS NO.: 199900061762 Reduction in HTLV-I proviral load and spontaneous lymphoproliferation in HTLV-I-associated myelopathy/tropical spastic paraparesis patients treated with humanized anti-Tac. AUTHOR: Lehky Tanya J(a); Levin Michael C; Kubota Ryuji; Bamford Richard N; Flerlage Alfred N; Soldan Samantha S; Leist Thomas P; Xavier Andrew; White Jeffrey D; Brown Margaret; Fleisher Thomas A; Top Lois E; Light Susan; McFarland Henry F; Waldmann Thomas A; Jacobson Steven AUTHOR ADDRESS: (a) HIV Res., DAIDS, NIAID, NIH, 6003 Executive Blvd., Rockville, MD 20892**USA JOURNAL: Annals of Neurology 44 (6):p942-947 Dec., 1998 ISSN: 0364-5134 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English 3/3/7 (Item 7 from file: 5) DIALOG(R) File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. BIOSIS NO.: 199799568066 Improved biodistribution of 125I-labeled anti-Tac disulfide-stabilized Fv fragment by blocking its binding to the alpha subunit of the interleukin 2 receptor in the circulation with preinjected humanized anti-Tac IgG. AUTHOR: Kobayashi Hisataka; Yoo Tae M; Drumm Debra; Kim Meyoung-Kon; Sun Bao-Fu; Le Nhat; Webber Keith O; Pastan Ira; Waldmann Thomas A; Paik Chang H; Carrasquillo Jorge A(a AUTHOR ADDRESS: (a) Nuclear Med. Dep., Build. 10, Room 1C496, NIH, 10 Center Drive, Bethesda, MD 20892-1180**USA JOURNAL: Cancer Research 57 (10):p1955-1961 1997 ISSN: 0008-5472 RECORD TYPE: Abstract LANGUAGE: English (Item 8 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. BIOSIS NO.: 199698609414 Radioimmunotherapy of interleukin-2R-alpha-expressing adult T-cell leukemia with Yttrium-90-labeled anti-tac. AUTHOR: Waldmann Thomas A(a); White Jeffrey D; Carrasquillo Jorge A; Reynolds James C; Paik Chang H; Gansow Otto A; Brechbiel Martin W; Jaffe

Elaine S; Fleisher Thomas A; Goldman Carolyn K; Top Lois E; Bamford Richard; Zaknoen Sara; Roessler Eric; Kasten-Sportes Claude; England Richard; Litou Hariklia; Johnson John A; Jackson-White Terri; Manns Angela; Hanchard Barrie; Junghans Richard P; Nelson David L AUTHOR ADDRESS: (a) Metabolism Branch, Natl. Cancer Inst. Health, Build. 10, Room 4N115, Bethesda, MD 20892**USA JOURNAL: Blood 86 (11):p4063-4075 1995 ISSN: 0006-4971 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English 3/3/9 (Item 9 from file: 5) DIALOG(R) File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. 10119190 BIOSIS NO.: 199698574108 The treatment of HAM/TSP patients with humanized anti-Tac antibody: Preliminary immunologic studies. AUTHOR: Lehky Tanya J; Levin Michael; Flerlage Nick; White Jeffrey; Fleisher Thomas; McFarland Henry; Waldmann Thomas; Jacobson Steven AUTHOR ADDRESS: NIH, Bethesda, MD**USA JOURNAL: Journal of Acquired Immune Deficiency Syndromes and Human Retrovirology 10 (2):p285 1995 CONFERENCE/MEETING: Seventh International Conference on Human Retrovirology HTLV and Related Viruses Paris, France October 17-21, 1995 ISSN: 1077-9450 RECORD TYPE: Citation LANGUAGE: English (Item 10 from file: 5) 3/3/10 DIALOG(R) File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. 09775306 BIOSIS NO.: 199598230224 Phenotypic knockout of the high-affinity human interleukin 2 receptor by intracellular single-chain antibodies against the alpha subunit of the AUTHOR: Richardson Jennifer H; Sodroski Joseph G; Waldmann Thomas A; Marasco Wayne A(a AUTHOR ADDRESS: (a) Dep. Med. Human Retrovirol., Dana-Farber Cancer Inst., Harvard Med. Sch., Boston, MA 02115**USA JOURNAL: Proceedings of the National Academy of Sciences of the United States of America 92 (8):p3137-3141 1995 ISSN: 0027-8424 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English 3/3/11 (Item 11 from file: 5) DIALOG(R) File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. BIOSIS NO.: 199598010657 09555739 Humanized Antibody Directed to the IL-2 Receptor beta-Chain Prolongs Primate Cardiac Allograft Survival. AUTHOR: Tinubu Sikiru A; Hakimi John; Kondas Jo A; Bailon Pascal; Familletti Philip C; Spence Cheryl; Crittenden Michael D; Parenteau Gary L; Dirbas Fredrick M; Tsudo Mitsuro; Bacher John D; Kasten-Sportes Claude ; Martinucci Jean L; Goldman Carolyn K; Clark Richard E; Waldmann Thomas A AUTHOR ADDRESS: Inq.**USA JOURNAL: Journal of Immunology 153 (9):p4330-4338 1994

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ISSN: 0022-1767 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English 3/3/12 (Item 12 from file: 5) DIALOG(R) File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. BIOSIS NO.: 199497463114 09454744 Treatment of acute graft-versus-host disease with humanized anti-gac: An antibody that binds to the interleukin-2 receptor. AUTHOR: Anasetti Claudio(a); Hansen John A; Waldmann Thomas A; Appelbaum Frederick R; Davis Jennifer; Deeg H Joachim; Doney Kristine; Martin Paul J; Nash Richard; et al AUTHOR ADDRESS: (a) Fred Hutchinson Cancer Res. Cent., 1124 Columbia St., E611, Seattle, WA 98104**USA JOURNAL: Blood 84 (4):p1320-1327 1994 ISSN: 0006-4971 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English (Item 13 from file: 5) DIALOG(R) File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. 09443625 BIOSIS NO.: 199497451995 Radioimmunotherapy of nude mice bearing a human interleukin 2 receptor alpha-expressing lymphoma utilizing the alpha-emitting radionuclide-conjugated monoclonal antibody 212Bi-anti-Tac. AUTHOR: Hartmann Frank; Horak Eva M; Garmestani Kayhan; Wu Chuanchu; Brechbiel Martin W; Kozak Robert W; Tso J; Kosteiny Sheri A; Gansow Otto A; Nelson David L; Waldmann Thomas A(a AUTHOR ADDRESS: (a) Metabolic Branch, Natl. Cancer Inst., Build. 10, 4N115, NIH, Bethesda, MD 20892**USA JOURNAL: Cancer Research 54 (16):p4362-4370 1994 ISSN: 0008-5472 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English (Item 14 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. BIOSIS NO.: 199497270183 Cooperative interactions between the interleukin 2 receptor alpha and beta chains alter the interleukin 2-binding affinity of the receptor subunits. AUTHOR: Roessler Erich(a); Grant Angus(a); Ju Grace; Tsudo Mitsuru; Sugamura Kazuo; Waldmann Thomas A(a AUTHOR ADDRESS: (a) Metabolism Branch, Natl. Cancer Inst., Natl. Inst. Health, Bethesda, MD 20892**USA

JOURNAL: Proceedings of the National Academy of Sciences of the United States of America 91 (8):p3344-3347 1994
ISSN: 0027-8424
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English

3/3/15 (Item 15 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)

(c) 2000 BIOSIS. All rts. reserv. 09039688 BIOSIS NO.: 199497048058 1992 Stohlman Memorial Lecture: Targeting the IL-2 receptor. AUTHOR: Waldmann Thomas A AUTHOR ADDRESS: Metabolism Branch, Natl. Cancer Inst., Natl. Inst. Health, Bethesda, MD 20892**USA JOURNAL: Leukemia (Basingstoke) 7 (SUPPL. 2):pS151-S156 1993 ISSN: 0887-6924 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English (Item 16 from file: 5) 3/3/16 DIALOG(R)File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. 08974421 BIOSIS NO.: 199396125922 The interleukin-2 receptor: A target for monoclonal antibody treatment of human T-cell lymphotropic virus I-induced adult T-cell leukemia. AUTHOR: Waldmann Thomas A(a); White Jeffrey D; Goldman Carolyn K; Top Lois; Grant Angus; Bamford Richard; Roessler Eric; Horak Ivan D; Zaknoen AUTHOR ADDRESS: (a) Metabolism Branch, Natl. Cancer Inst., Build. 4N115, Bethesda, MD 20892**USA JOURNAL: Blood 82 (6):p1701-1712 1993 ISSN: 0006-4971 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English (Item 17 from file: 5) 3/3/17 DIALOG(R)File 5:Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. BIOSIS NO.: 199396042247 08890746 Cytotoxic activities of recombinant immunotoxins composed of Pseudomonas toxin or diphtheria toxin toward lymphocytes from patients with adult T-cell leukemia. AUTHOR: Kreitman Robert J; Chaudhary Vijay K; Waldmann Thomas A; Hanchard Barrie; Cranston Beverly; Fitzgerald David J P; Pastan Ira(a AUTHOR ADDRESS: (a) Lab. Mol. Biol., Div. Cancer Biol., Diagnosis Cent., NCH NIH, 9000 Rockville Pike, 37/4E16, Beth JOURNAL: Leukemia (Basingstoke) 7 (4):p553-562 1993 ISSN: 0887-6924 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English 3/3/18 (Item 18 from file: 5) DIALOG(R) File 5: Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv. BIOSIS NO.: 199395074373 Prolongation of graft survival in primate allograft transplantation by yttrium-90-labeled anti-Tac in conjunction with granulocyte colony-stimulating factor.

AUTHOR: Parenteau Gary L; Dirbas Frederick M; Garmestani Kayhan; Brechbiel Martin W; Bukowski Maria A; Goldman Carolyn K; Clark Richard; Gansow Otto A; Waldmann Thomas A(a

AUTHOR ADDRESS: (a) Metabolism Branch, National Cancer Inst., Bldg. 10, Room 4N115, National Inst. Health, Bethesda,

JOURNAL: Transplantation (Baltimore) 54 (6):p963-968 1992

, .ISSN: 0041-1337

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

3/3/19 (Item 19 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
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08751679 BIOSIS NO.: 199395041030

Recombinant toxins containing the variable domains of the anti-Tac monoclonal antibody to the interleukin-2 receptor kill malignant cells from patients with chronic lymphocytic leukemia.

AUTHOR: Kreitman Robert J; Chaudhary Vijay K; Kozak Robert W; Fitzgerald David J P; Waldmann Thomas A; Pastan Ira(a

AUTHOR ADDRESS: (a) Lab. Mol. Biol., Div. Cancer Biol., Diagnosis and

Centers, Natl. Cancer Inst., Natl. Inst. Healt

JOURNAL: Blood 80 (9):p2344-2352 1992

ISSN: 0006-4971

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

3/3/20 (Item 20 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2000 BIOSIS. All rts. reserv.

08749929 BIOSIS NO.: 199395039280

Mik-beta-1(Fv)-PE40, a recombinant immunotoxin cytotoxic toward cells bearing the beta-chain of the IL-2 receptor.

AUTHOR: Kreitman Robert J; Schneider William P; Queen Cary; Tsudo Mitsuru;

Fitzgerald David J P; Waldmann Thomas A; Pastan Ira(a

AUTHOR ADDRESS: (a) Lab. Mol. Biol./Div. Cancer Biol., Diagnosis and

Centers, 37/4E16, Natl. Cancer Inst., Natl. Ins

JOURNAL: Journal of Immunology 149 (8):p2810-2815 1992

ISSN: 0022-1767

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English potential applications in organ transplantation and the **treatment** of IL-2 receptor-expressing neoplastic diseases.

2/7/3 (Item 3 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
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07421848 BIOSIS NO.: 000091027837 USE OF YTTRIUM-90-LABELED ANTI-TAC ANTIBODY IN PRIMATE XENOGRAFT TRANSPLANTATION

AUTHOR: COOPER M M; ROBBINS R C; GOLDMAN C K; MIRZADEH S; BRECHBIEL M W; STONE C D; GANSOW O A; CLARK R E; WALDMANN T A

AUTHOR ADDRESS: DEP. OF SURG., COLUMBIA UNIV. COLL. OF PHYSICIANS AND SURGEONS, 630 WEST 168TH ST., NEW YORK, N.Y. 10032.

JOURNAL: TRANSPLANTATION (BALTIMORE) 50 (5). 1990. 760-765.

FULL JOURNAL NAME: TRANSPLANTATION (Baltimore)

CODEN: TRPLA

RECORD TYPE: Abstract LANGUAGE: ENGLISH

ABSTRACT: The high-affinity interleukin-2 receptor (IL-2R) is expressed by T cell activated in response to foreign histocompatibility antigens but not by normal resting cells. Thus, blockade of the interaction of IL-2 with its receptor could achieve selective immunosuppression. Accordingly, anti-Tac, a murine IgG2a class monoclonal antibody specific to the IL-2R, was used alone or in a chelated form with yttrium-90 (90Y), a pure beta emitter, to inhibit rejection of cardiac xenografts from Macaca fascicularis (cynomolgus) donors transplanted to the cervical or abdominal region of Macaca mulatta (rhesus) recipients (n = 20). Animals received no immunosuppression (n = 3, group I, controls), unmodified anti-Tac (n = 5, 2 mg/kg q.o.d., group II), or 90Y-anti-Tac (n = 5, 16 mCi, group III). To distinguish the nonspecific immunosuppressive effect of radiation, 90Y was administered bound to UPC-10 (n = 4, 16 mCi, group IV), another murine monoclonal antibody that does not specifically recognize activated immunoresponsive cells. All immunosuppression was administered in divided doses during the first 2 weeks posttransplant. Group I animals rejected their grafts at 6.7 .+-. 1 days and demonstrated a rise is soluble IL-2R levels at the time of rejection, indicating the generation of Tac-expressing and -releasing cells. Graft survival in group II was not prolonged compared with controls (mean survival 6.2 .+-. 1 days; P > 0.05). In contrast, graft survival in animals that received the designed dosage of 90Y -anti-Tac was significantly prolonged to an average of 38.4 .+-. 5 days compared with groups I and II (P < 0.005 and P < 0.005, respectively). Prolongation of graft survival occurred in animals that received 90Y-UPC-10 (mean survival 21.3 .+-. 5 days, P < 0.05 versus group I, P < 0.01 versus group II). However, 90Y-UPC-10 was significantly less effective in prolonging graft survival than 90Y -anti-Tac, in which one-half the per-kilogram dosage of radioactivity was delivered in specific fashion via anti-Tac (P < 0.025). Reversible nonlethal bone marrow suppression occurred without associated nephro- or hepatotoxicity, and virtually all animals developed antibodies to the urine monoclonal. Thus, the approach used in the present study, IL-2R-directed therapy with 90Y-anti-Tac, may have potential applications in organ transplantation and in the treatment of Tac-expressing neoplastic diseases.

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$0.46 Estimated total session cost 0.173 DialUnits
File 357: Derwent Biotechnology Abs 1982-2000/Jun B1
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       Set Items Description
? e au=waldmann thomas
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E1 1 AU=WALDMANN R; +LOCKSLEY R M; GLAICHENHAUS N
E2 9 AU=WALDMANN T A
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E3 0 *AU=WALDMANN THOMAS
E4 1 AU=WALDMANN V
E5 1 AU=WALDMEIER P
E6 1 AU=WALDNER H
E7 10 AU=WALDNER R
E8 2 AU=WALDNER-SANDER S
E9 2 AU=WALDO G S
E10 1 AU=WALDO III R H
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E12 5 AU=WALDRAFF W
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DIALOG(R) File 357: Derwent Biotechnology Abs
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0234528 DBA Accession No.: 99-04629
                                            PATENT
Killing cells with immunotoxin - cell lysis using a Pseudomonas sp.
     exotoxin and single chain antibody scFv fragment fusion, antibody
     engineering for autoimmune disease and cancer therapy
AUTHOR: Fitzgerald D J; Chaudhary V K; Pastan I H; Waldmann T A;
CORPORATE SOURCE: Washington, DC, USA.
PATENT ASSIGNEE: U.S.Dep.Health-Hum.Serv. 1999
PATENT NUMBER: US 5863745 PATENT DATE: 990126 WPI ACCESSION NO.:
     99-131300 (9911)
PRIORITY APPLIC. NO.: US 461825 APPLIC. DATE: 950605
NATIONAL APPLIC. NO.: US 461825 APPLIC. DATE: 950605
LANGUAGE: English
DIALOG(R) File 357: Derwent Biotechnology Abs
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0220988 DBA Accession No.: 98-02585
                                             PATENT
Fusion protein of antibody Fv fragment and Pseudomonas exotoxin fragment -
    an immunotoxin for use as e.g. an antitumor or immunosuppressive agent
AUTHOR: FitzGerald D; Chaudhary V K; Pastan I H; Waldmann T A;
    Queen C L
CORPORATE SOURCE: Washington, DC, USA.
PATENT ASSIGNEE: U.S.Dep.Health-Hum.Serv. 1997
PATENT NUMBER: US 5696237 PATENT DATE: 971209 WPI ACCESSION NO.:
    98-041352 (9804)
PRIORITY APPLIC. NO.: US 463163 APPLIC. DATE: 950605
NATIONAL APPLIC. NO.: US 463163 APPLIC. DATE: 950605
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\$0.00 Estimated cost this search

LANGUAGE: English

1/3/3 DIALOG(R)File 357:Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv. 0164255 DBA Accession No.: 94-06806 Genetically engineered monoclonal antibodies armed with radionuclides -

radionuclide-carrying anti-Tac monoclonal antibody with reduced immunogenicity (humanized antibody engineering) for interleukin-2 receptor-directed therapy (conference paper) AUTHOR: Waldmann T A

CORPORATE AFFILIATE: Nat.Cancer-Inst.Bethesda Nat.Inst.Health-Bethesda CORPORATE SOURCE: Metabolism Branch, Bldg 10, Rm 4N115, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892, USA.

JOURNAL: BIOTECH'92 (7, 205-12) 1993

CODEN: 9999Q LANGUAGE: English

1/3/4

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0123619 DBA Accession No.: 91-11261

Monoclonal antibodies in diagnosis and therapy - monoclonal antibody production; a review

AUTHOR: Waldmann T A

CORPORATE SOURCE: Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892, USA.

JOURNAL: Science (252, 5013, 1657-62) 1991

CODEN: SCIEAS LANGUAGE: English

1/3/5

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0110818 DBA Accession No.: 90-13509

IL2-PE664Glu, a new chimeric protein cytotoxic to human-activated T-lymphocytes - human interleukin-2 gene fusion with Pseudomonas exotoxin mutant gene; potential use of chimeric toxin as cytostatic and immunosuppressive

AUTHOR: Loberboum-Galski H; Garsia R J; Gately M; Brown P S; Clark R E; Waldmann T A

CORPORATE AFFILIATE: Roche

CORPORATE SOURCE: Laboratory of Molecular Biology, Division of Cancer Biology and Diagnosis, National Cancer Institute, National Institutes of Health, Bethesda, Maryland 20892, USA.

JOURNAL: J.Biol.Chem. (265, 27, 16311-17) 1990

CODEN: JBCHA3 LANGUAGE: English

1/3/6

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0090949 DBA Accession No.: 89-08940

A recombinant immunotoxin consisting of two antibody variable domains fused to Pseudomonas exotoxin - monoclonal antibody anti-Tac to the p55 subunit of human interleukin-2 receptor

AUTHOR: Chaudhary V K; Queen C; Junghans R P; Waldmann T A; FitzGerald D J; Pastan I

CORPORATE AFFILIATE: Protein-Design-Labs

CORPORATE SOURCE: Laboratory of Molecular Biology, DCBD, National Cancer Institute, National Institutes of Health, Bethesda, Maryland 20892,

JOURNAL: Nature (339, 6223, 394-97) 1989

CODEN: NATUAS LANGUAGE: English

1/3/7

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0054663 DBA Accession No.: 86-12511

Radionuclide-conjugated monoclonal antibodies: a synthesis of immunology, inorganic chemistry and nuclear science - development and application in tumor diagnosis etc.

AUTHOR: Kozak R W; Waldmann T A; Atcher R W; Gansow O A CORPORATE SOURCE: Metabolism Branch, National Cancer Institute of NIH, Bethesda, MD 20892, USA.

JOURNAL: Trends Biotechnol. (4, 10, 259-64) 1986

CODEN: 8921M LANGUAGE: English

1/3/8

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0049478 DBA Accession No.: 86-07326

The structure, function, and expression of interleukin-2 receptors on normal and malignant lymphocytes - investigated using a monoclonal antibody

AUTHOR: Waldmann T A

CORPORATE SOURCE: Metabolism Branch, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892, USA.

JOURNAL: Science (232, 4751, 727-32) 1986

CODEN: SCIEAS LANGUAGE: English

1/3/9

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0005909 DBA Accession No.: 82-04909

A monoclonal antibody that appears to recognize the receptor for human T-cell growth factor; partial characterization of the receptor - purification of the interleukin-2 surface receptor by immunoprecipitation

AUTHOR: Leonard W J; Depper J M; Uchiyama T; Smith K A; Waldmann T A; Greene W C

CORPORATE SOURCE: The Metabolism Branch, National Cancer Institute, NIH, Bethesda, Maryland 20205, USA.

JOURNAL: Nature (300, 5889, 267-69) 1982

CODEN: NATUAS LANGUAGE: English DIALOG(R)File 357:Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv. 0234528 DBA Accession No.: 99-04629 PATENT Killing cells with immunotoxin - cell lysis using a Pseudomonas sp. exotoxin and single chain antibody scFv fragment fusion, antibody engineering for autoimmune disease and cancer therapy AUTHOR: Fitzgerald D J; Chaudhary V K; Pastan I H; Waldmann T A; Queen C L CORPORATE SOURCE: Washington, DC, USA. 1999 PATENT ASSIGNEE: U.S.Dep.Health-Hum.Serv. PATENT NUMBER: US 5863745 PATENT DATE: 990126 WPI ACCESSION NO.: 99-131300 (9911) PRIORITY APPLIC. NO.: US 461825 APPLIC. DATE: 950605 NATIONAL APPLIC. NO.: US 461825 APPLIC. DATE: 950605 LANGUAGE: English 1/3/2 DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv. 0220988 DBA Accession No.: 98-02585 PATENT Fusion protein of antibody Fv fragment and Pseudomonas exotoxin fragment an immunotoxin for use as e.g. an antitumor or immunosuppressive agent AUTHOR: FitzGerald D; Chaudhary V K; Pastan I H; Waldmann T A; Queen C L CORPORATE SOURCE: Washington, DC, USA. PATENT ASSIGNEE: U.S.Dep.Health-Hum.Serv. 1997 PATENT NUMBER: US 5696237 PATENT DATE: 971209 WPI ACCESSION NO.: 98-041352 (9804) PRIORITY APPLIC. NO.: US 463163 APPLIC. DATE: 950605 NATIONAL APPLIC. NO.: US 463163 APPLIC. DATE: 950605 LANGUAGE: English 1/3/3 DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv. 0164255 DBA Accession No.: 94-06806 Genetically engineered monoclonal antibodies armed with radionuclides radionuclide-carrying anti-Tac monoclonal antibody with reduced immunogenicity (humanized antibody engineering) for interleukin-2 receptor-directed therapy (conference paper) AUTHOR: Waldmann T A CORPORATE AFFILIATE: Nat.Cancer-Inst.Bethesda Nat.Inst.Health-Bethesda CORPORATE SOURCE: Metabolism Branch, Bldg 10, Rm 4N115, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892, USA. JOURNAL: BIOTECH'92 (7, 205-12) 1993 CODEN: 9999Q LANGUAGE: English DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0123619 DBA Accession No.: 91-11261

Monoclonal antibodies in diagnosis and therapy - monoclonal antibody production; a review

AUTHOR: Waldmann T A

CORPORATE SOURCE: Metabolism Branch, National Cancer Institute, National

Institutes of Health, Bethesda, MD 20892, USA. JOURNAL: Science (252, 5013, 1657-62) 1991

CODEN: SCIEAS LANGUAGE: English

1/3/5

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0110818 DBA Accession No.: 90-13509

IL2-PE664Glu, a new chimeric protein cytotoxic to human-activated
 T-lymphocytes - human interleukin-2 gene fusion with Pseudomonas
 exotoxin mutant gene; potential use of chimeric toxin as cytostatic and
 immunosuppressive

AUTHOR: Loberboum-Galski H; Garsia R J; Gately M; Brown P S; Clark R E; Waldmann T A

CORPORATE AFFILIATE: Roche

CORPORATE SOURCE: Laboratory of Molecular Biology, Division of Cancer Biology and Diagnosis, National Cancer Institute, National Institutes of Health, Bethesda, Maryland 20892, USA.

JOURNAL: J.Biol.Chem. (265, 27, 16311-17) 1990

CODEN: JBCHA3 LANGUAGE: English

1/3/6

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0090949 DBA Accession No.: 89-08940

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AUTHOR: Chaudhary V K; Queen C; Junghans R P; Waldmann T A; FitzGerald D J; Pastan I

CORPORATE AFFILIATE: Protein-Design-Labs

CORPORATE SOURCE: Laboratory of Molecular Biology, DCBD, National Cancer Institute, National Institutes of Health, Bethesda, Maryland 20892, USA.

JOURNAL: Nature (339, 6223, 394-97) 1989

CODEN: NATUAS LANGUAGE: English

1/3/7

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0054663 DBA Accession No.: 86-12511

Radionuclide-conjugated monoclonal antibodies: a synthesis of immunology, inorganic chemistry and nuclear science - development and application in tumor diagnosis etc.

AUTHOR: Kozak R W; Waldmann T A; Atcher R W; Gansow O A

CORPORATE SOURCE: Metabolism Branch, National Cancer Institute of NIH, Bethesda, MD 20892, USA.

JOURNAL: Trends Biotechnol. (4, 10, 259-64) 1986

CODEN: 8921M LANGUAGE: English DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0049478 DBA Accession No.: 86-07326

The structure, function, and expression of interleukin-2 receptors on normal and malignant lymphocytes - investigated using a monoclonal antibody

AUTHOR: Waldmann T A

CORPORATE SOURCE: Metabolism Branch, National Cancer Institute, National

Institutes of Health, Bethesda, MD 20892, USA.

JOURNAL: Science (232, 4751, 727-32) 1986

CODEN: SCIEAS LANGUAGE: English

1/3/9

DIALOG(R) File 357: Derwent Biotechnology Abs (c) 2000 Derwent Publ Ltd. All rts. reserv.

0005909 DBA Accession No.: 82-04909

A monoclonal antibody that appears to recognize the receptor for human T-cell growth factor; partial characterization of the receptor - purification of the interleukin-2 surface receptor by immunoprecipitation

AUTHOR: Leonard W J; Depper J M; Uchiyama T; Smith K A; Waldmann T A; Greene W C

CORPORATE SOURCE: The Metabolism Branch, National Cancer Institute, NIH, Bethesda, Maryland 20205, USA.

JOURNAL: Nature (300, 5889, 267-69) 1982

CODEN: NATUAS LANGUAGE: English

(Item 13 from file: 73) DIALOG(R) File 73: EMBASE

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EMBASE No: 1998050169

Clinical pharmacology and therapeutic potential of monoclonal antibody treatment in rheumatoid arthritis

Choy E.H.S.

Dr. E.H.S. Choy, Rheumatology Unit, Thomas Guy House, Guy's Hospital, St Thomas Street, London SE1 9RT United Kingdom

AUTHOR EMAIL: e.choy@umds.ac.uk

Drugs and Aging (DRUGS AGING) (New Zealand) 1998, 12/2 (139-148)

CODEN: DRAGE ISSN: 1170-229X DOCUMENT TYPE: Journal; Review

SUMMARY LANGUAGE: ENGLISH LANGUAGE: ENGLISH

7/3/38 (Item 9 from file: 73)
DIALOG(R)File 73:EMBASE
(c) 2000 Elsevier Science B.V. All rts. reserv.

07247945 EMBASE No: 1998138658

Biological agents in rheumatoid arthritis: Which ones could be used in combination?

Lorenz H.-M.; Kalden J.R.

Dr. H.-M. Lorenz, Department of Medicine III, Inst. Clinical Immunol./Rheumatology, University of Erlangen-Nuremberg,

Krankenhausstrasse 12, 91054 Erlangen Germany

AUTHOR EMAIL: Hannes.Lorenz@med3.med.uni-erlangen.de BioDrugs (BIODRUGS) (New Zealand) 1998, 9/4 (303-324)

CODEN: BIDRF ISSN: 1173-8804 DOCUMENT TYPE: Journal; Review

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 130

7/3/32 (Item 3 from file: 73)

DIALOG(R) File 73: EMBASE

(c) 2000 Elsevier Science B.V. All rts. reserv.

07932003 EMBASE No: 1999405331

Novel biologic approaches to the treatment of rheumatoid $\operatorname{arthritis}$ Sany J.

Prof. J. Sany, Service d'Immuno-Rhumatologie, Hopital Lapeyronie, 34295 Montpellier Cedex 2 France

Revue du Rhumatisme (English Edition) (REV. RHUM. ENGL. ED.) (France)

1999, 66/11 (548-559)

CODEN: RRHUE ISSN: 1169-8446

DOCUMENT TYPE: Journal; Conference Paper

LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 73

7/3/33 (Item 4 from file: 73)

DIALOG(R) File 73: EMBASE

(c) 2000 Elsevier Science B.V. All rts. reserv.

07768530 EMBASE No: 1999247338

What we have learned from trials of immunomodulatory agents in rheumatoid arthritis: Future directions

Schulze-Koops H.; Burkhardt H.; Kalden J.R.

Dr. H. Schulze-Koops, Department of Internal Medicine III, University of Erlangen-Nuremberg, Krankenhausstrasse 12, D-91054 Erlangen Germany

Drugs of Today (DRUGS TODAY) (Spain) 1999, 35/4-5 (327-351)

CODEN: MDACA ISSN: 0025-7656

DOCUMENT TYPE: Journal; Conference Paper

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 172

(Item 28 from file: 5) DIALOG(R) File 5: Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv.

07982165 BIOSIS NO.: 000042033563 HUMANIZED MONOCLONAL ANTIBODY TREATMENT IN RHEUMATOID ARTHRITIS AUTHOR: KYLE V; RODDY J; HALE G; HAZLEMAN B L; WALDMANN H AUTHOR ADDRESS: RHEUMATOLOGY RESEARCH UNIT, ADDENBROOKE'S HOSPITAL,

CAMBRIDGE, UK.

JOURNAL: J RHEUMATOL 18 (11). 1991. 1737-1738. FULL JOURNAL NAME: Journal of Rheumatology

CODEN: JRHUA

RECORD TYPE: Citation LANGUAGE: ENGLISH

(Item 29 from file: 5) 7/3/29 DIALOG(R) File 5: Biosis Previews(R) (c) 2000 BIOSIS. All rts. reserv.

06744219 BIOSIS NO.: 000088053649 BENEFICIAL EFFECT OF MONOCLONAL ANTIBODY TO INTERLEUKIN 2 RECEPTOR ON ACTIVATED T CELLS IN RHEUMATOID ARTHRITIS AUTHOR: KYLE V; COUGHLAN R J; TIGHE H; WALDMANN H; HAZLEMAN B L AUTHOR ADDRESS: RHEUMATOL. RES. UNIT., UNIT E6, ADDENBROOKE'S HOSP., HILLS

ROAD, CAMBRIDGE CB2 2QQ.

JOURNAL: ANN RHEUM DIS 48 (5). 1989. 428-429.

FULL JOURNAL NAME: Annals of the Rheumatic Diseases

CODEN: ARDIA

RECORD TYPE: Abstract